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# COMPOSITIONS AND METHODS FOR TREATING OR PREVENTING HIV INFECTION

This application claims the benefit of U.S. Provisional Application Nos. 60/491,258 filed July 31, 2003, 60/493,767 filed August 11, 2003, 60/496,908 filed August 22, 2003, and 60/501,832 filed September 11, 2003, which are hereby incorporated by reference in their entirety.

### BACKGROUND OF THE INVENTION

Acquired Immune Deficiency Syndrome ("AIDS") is one of the most serious health threats confronting the human population today. AIDS is caused by a virus known as human immunodeficiency virus ("HIV") which presently includes HIV-1 and HIV-2. Over 40 million people are estimated to be living with HIV/AIDS. Current projections suggest that an additional 45 million people will become infected between 2002 and 2010. So far, it is believed that more 25 million people have died from AIDS.

Since its emergence in the 1970s, HIV has produced a continually growing global pandemic, and it has, thus far, defied all attempts to produce an effective vaccine. Although a number of drugs have been developed to treat the disease, all have limited usefulness, serious side effects, a high potential for resistance, and none have been identified so far which can cure or prevent it. HIV vaccine research has expanded over recent years, but success so far using HIV-based components has been limited. See, e.g., Graham et al., *J. Inf. Disease.*, 166:244-252, 1992; Belshe et al., *J. Inf. Disease.*, 183:1343-52, 2001; Horton et al., *J. Virol.*, 76:7187-7202, 2002; Gilbert et al., *Vaccine*, 21:2933-2947, 2003.

#### **DESCRIPTION OF DRAWINGS**

FIG. 1 (A-C). Comparison of cells from vaccinated versus non-vaccinated subjects, infected with the macrophage (CCR5) tropic HIV. A. A comparison of the mean + standard error measurement of the vaccinated versus non-vaccinated groups in cultures without autologous serum. (\*, p<0.05) B. A comparison of the mean +

standard error measurement of the vaccinated versus non-vaccinated groups in cultures with autologous serum (\*, p< 0.05; \*\*, p<0.01). C. Comparison of the mean + standard error measurement of cells from vaccinated versus non-vaccinated subjects, infected with the T-cell (CXCR4) tropic HIV.

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### DESCRIPTION OF THE INVENTION

The present invention provides methods and compositions for treating and/or preventing HIV infection in a subject in need thereof. It features the use of poxviruses for therapy, prophylaxis, and diagnosis of HIV, as well as for any other medical or veterinary use associated with HIV and homologous viruses. The invention also provides for the use of poxviruses in the discovery of new agents to prevent and/or treat HIV infection.

A poxvirus or a component thereof, can be used to treat and/or prevent infection caused by any virus, preferably a lentivirus, such as HIV, that uses a CCR5 chemokine receptor for its infection of cells. This includes, but is not limited to, e.g., HIV-1 (e.g., clades A, B, C, D, and G, R5 and R5X4 viruses, etc.), HIV-2 (e.g., R5 and R5X4 viruses, etc.), simian immunodeficiency virus (SIV), simian/human immunodeficiency virus (SHIV), feline immunodeficiency virus (FIV), bovine immunodeficiency virus (BIV) (Wright et al., *Vet. Res. Commun.*, 26:239-50, 2002), HTLV-1, HTLV-2, etc. It can be used as a vaccine, adjuvant, therapeutic agent, in combination with other agents, or in any suitable manner to treat and/or prevent such infections.

Any poxvirus can be used in accordance with the present invention, including, but not limited to, orthopoxvirus, parapoxvirus, avipoxvirus, capripoxvirus, leporipoxvirus, suipoxvirus, etc. Orthopoxvirus, include, e.g., buffalopox, camelpox, cowpox, monkeypox, rabbitpox, raccoon pox, tatera pox, canarypox, fowlpox, vaccinia, variola, and vole pox. Vaccinia virus is the prototype of the genus Orthopoxvirus for the desired effects, but other poxviruses can be used in its place. Thus, although the disclosure below may be written in terms of vaccinia, any poxvirus can be utilized in accordance with the present invention.

Vaccinia is a double-stranded DNA (deoxyribonucleic acid) virus. All strains, derivatives, variants, mutations, naturally-occurring strains, genetically-engineered, recombinant, etc., of vaccinia can be used in accordance with the present invention. For more information on vaccinia and other poxvirus, see e.g., *Virology*, Fields et al., Volume 2, Chapters 74-75, Raven Press, 1990.

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An amount of the poxvirus, such as vaccinia virus, can be administered to a subject in a quantity which is effective to achieve a therapeutic or prophylactic effect. The term "poxvirus," "vaccinia virus," etc., indicates that the virus (genome and protein coat) is administered to a subject. It can be administered in any effective form, including, e.g., as a live virus, as a live-attenuated virus, as a replication-deficient virus, as a viral extract not having any live viral particles, etc. Compositions comprising a poxvirus can be produced and utilized in any suitable manner, including, e.g., recombinant, naked DNA technology, etc.

The term "treating" is used conventionally, e.g., the management or care of a subject for the purpose of combating, alleviating, reducing, relieving, improving, eliminating, etc., one or more signs or symptoms associated with HIV infection.

Treatment includes delaying the progression of HIV and its associated symptoms, thereby extending the life expectancy of an infected subject, and/or delaying or reducing the onset of symptoms associated with HIV infection. Treating can involve inhibiting, reducing, diminishing, etc., the replication and other events in the life cycle of the HIV virus.

The term "preventing" HIV infection indicates that a subject's susceptibility to HIV infection upon exposure to the virus is reduced or diminished as a result of the administration of the poxvirus. The subject's resistance to HIV infection is increased or improved by the poxvirus treatment since s/he is less likely to become infected by the virus. Any amount of improved resistance is useful, e.g., greater than 5-fold, greater than 7-fold, greater than ten-fold, etc., and any such improvement can be regarded as prevention.

A poxvirus, or component thereof, used in the present invention can be prepared routinely, or obtained from commercial sources. Attenuated strains are preferred. Attenuated strains are less able to cause disease, and are considered less virulent and weakened as compared to strains that are not attenuated.

Any strain of vaccinia virus, or components thereof, can be utilized to achieve a prophylactic and/or therapeutic effect, including, but not limited to, e.g., strains available from the ATCC, ECACC, or other virus collections, replication-competent. replication-deficient, non-replicating, attenuated strains, modified vaccinia Ankara (MVA), vaccinia virus Ankara, NYVAC (ATCC No. VR-2559) replication-deficient vaccinia viruses, VV Copenhagen, VV Western Reserve, VV Wyeth (ATCC No. VR325), Elstree, strains deficient in vCCI (Reading et al., J. Immunol., 170:1435-42, 2003), and/or vGF, strains comprising one or more copies of the 17K myristyloprotein, poxvirus strains, CCR5-dependent poxvirus strains, etc. Dryvax®, 10 a vaccinia (smallpox) vaccine currently licensed in the United States, is a lyophilized, live-virus preparation of infectious vaccinia virus (Wyeth Laboratories, Inc., Marietta, Pennsylvania). Other strains which have been used include, but are not limited to. e.g., Lister, Bordeaux, Paris, Massachusetts 999, New York, Temple of Heaven, Patwadangar, Ikeda, Bern, Vienna, Bohemia, Finland, Hamburg, Budapest, Aosta, Spain, Sweden, B-51, Tashkent, EM-63, LE-IVP (Lister), etc. See, also, Smallpox and 15 its Eradication, Fenner et al., WHO, Geneva, 1988, e.g., Chapter 11. Other strains include, e.g., MVA-BN (modified vaccinia Ankara - Bavarian Nordic) (ECACC V00083008; WO 02/42480), MVA-Vero (US 20030013190), MVA-NH, MVA 572 (ECACC V94012707), LC16m8, and ACAM1000 (ATCC Deposit No. PTA-3321; 20 WO 02/085411). Any strain of canarypox can be utilized as well, including attenuated canarypox virus such as, e.g., ALVAC (ATCC No. VR-2547). Deposited strains also include, e.g., ATCC Nos. VR-117 (CL), VR-118 (Lederle-Chorioallantoic), VR-119 (WR (Mouse Neurotropic), VR-1354 (WR (NIH TC-adapted), VR-1431 (P-4), VR-1441 (IHD-W), VR-1508 (Modified vaccinia virus

Deposited strains also include, e.g., ATCC Nos. VR-117 (CL), VR-118 (Lederle-Chorioallantoic), VR-119 (WR (Mouse Neurotropic), VR-1354 (WR (NIH TC-adapted), VR-1431 (P-4), VR-1441 (IHD-W), VR-1508 (Modified vaccinia virus Ankara (MVA)), VR-1536 (New York City Department of Health Laboratories (Wyeth-calf adapted)), VR-1549 (Elstree (Lister Vaccine)), VR-156 (IHD), VR-2010 (AS), VR-2031 (Vtk-79), VR-2034 (S-variant), VR-2042 (vP-7), VR-2043 (vP-9), VR-2292 (SLZ103[recombinant Vaccinia virus (WR)]), VR-2379 (Rpmuhr+ [recombinant of Utrecht strain Rpuhr23]), VR-2589 (VVtm1:hPC1 [recombinant Vaccinia virus, in vitro construct]), VR-302 (Brighton), VR-3103 (IHD-W Dts 16 [Vaccinia ts-mutant]), VR-3109 (IHD-W Dts 46 [Vaccinia ts-mutant]), VR-3110 (IHD-W Dts 2 [Vaccinia ts-mutant]), VR-3113 (IHD-W Dts 17 [Vaccinia ts-mutant]),

VR-3121 (IHD-W Dts8 [Vaccinia ts-mutant]), VR-3126 (IHD-W Dts 33 [Vaccinia ts-mutant]), VR-3129 (IHD-W Dts 48 [Vaccinia ts-mutant]), VR-3130 (IHD-W Dts 4 [Vaccinia ts-mutant]), VR-3139 (IHD-W Dts 50 [Vaccinia ts-mutant]), VR-3142 (IHD-W Dts 10 [Vaccinia ts-mutant]), VR-3144 (IHD-W Dts20), VR-3147 (IHD-W Dts 35 [Vaccinia ts-mutant]), VR-3148 (IHD-W Dts 40), VR-3154 (IHD-W Dts71 [Vaccinia ts-mutant]), VR-3160 (IHD-W Dts52 [Vaccinia ts-mutant]), VR-3161 (IHD-W Dts 57), VR-3165 (IHD-W Dts 77), VR-3166 (IHD-W Dts 82), VR-3169 (IHD-W Dts97 [Vaccinia ts-mutant]), VR-3175 (IHD-W Dts 78 [Vaccinia ts-mutant]), VR-3176 (IHD-W Dts 83 [Vaccinia ts-mutant]), VR-3178 (IHD-W Dts 93 [Vaccinia ts-mutant]), VR-3196 (IHD-W Dts 95 [Vaccinia ts-mutant]), VR-587 (Yaba monkey tumor virus deposited as Yaba monkey tumor virus, Yatapoxvirus (Roswell Park-Yohn)), VR-838 (Raccoonpox virus, Orthopoxvirus (Herman)).

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A vaccinia virus is a preferred poxvirus in accordance with the present invention, but other poxviruses can also be used to treat and/or prevent HIV. For example, any poxvirus which expresses a gp120-like or TAT-like polypeptide, or which depends on CCR5 for entry into a cell can be used in accordance with the present invention.

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Vaccinia virus can be administered to subjects according to any regimen which is effective for treating and/or preventing HIV infection. The particular dosages (i.e., effective amounts), and number and frequency of vaccinations can be determined routinely.

An effective amount of virus, or virus component, is the quantity of virus, or virus component, which is useful to achieve the desired purpose, e.g., to treat and/or prevent HIV infection. These amounts can be determined routinely. Effective amounts can be the same or less than the amounts currently used to achieve pox immunity with a pox vaccine. For example, Dryvax<sup>TM</sup> is commonly used at a potency of 100 million pock-forming units (pfu)/ml for primary vaccination for smallpox. Any effective amount can be used in accordance with the present invention, e.g., about 10<sup>5</sup>-10<sup>9</sup> pfu/ml. The quantities of the particular virus which is utilized can be adjusted and determined routinely, e.g., to eliminate or reduce adverse reactions associated with the virus, as well as depending on the health of the patient receiving the treatment.

The specific dose level and frequency of dosage may vary, and can depend upon a variety of factors, including the activity and state of the specific poxvirus, e.g., whether it is live, heat-inactivated, attenuated, etc., its metabolic stability and length of action, rate of excretion, mode and time of administration, and the age, body weight, general health, gender, diet, and particular condition of the subject undergoing treatment or prevention.

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Poxvirus can be administered in any form by any effective route, including, e.g., oral, parenteral, enteral, intraperitoneal, topical, transdermal (e.g., using any standard patch), ophthalmic, nasally, local, non-oral, such as aerosal, spray, inhalation, percutaneous (epidermal), subcutaneous, intravenous, intramuscular, buccal, sublingual, rectal, vaginal, intra-arterial, mucosal, and intrathecal, etc. It can be administered alone, or in combination with any ingredient(s), active or inactive.

Any subject can be administered a poxvirus in accordance with the present invention, including subjects who have been exposed to HIV, but have not yet developed HIV infection, as well as subjects who have progressed to one or more of the clinical symptoms of HIV infection (e.g., AIDS). In addition to treating and/or preventing HIV infection in humans, a poxvirus can be used to treat other organisms (e.g., non-human primates, cats, etc.) infected with HIV, or HIV-related viruses, such as SIV, SHIV, or FIV. Thus, subjects who can be treated include, e.g., mammals, humans, monkeys, apes, chimpanzees, gorillas, cats, dogs, mice, rats, etc.

Subjects, who have been exposed to HIV virus, or who are at risk for developing the disease, are particular candidates for poxvirus vaccination. For instance, a subject who has not yet tested positive, but has been exposed to HIV, can be administered vaccinia virus as a prophylactic/therapeutic approach. Individuals at high-risk for the disease, such as sexually-active individuals, subjects in parts of the world where HIV infection is high, subjects receiving blood and/or other invasive medical procedures, can also receive vaccination to increase their resistance to HIV infection.

In addition to administering the whole poxvirus, components of it can also be administered in accordance with the present invention. By the phrase "component," it is meant any part of the virus, which is less than the whole virus genome, including

particular nucleic segments of its genome, as well as any product which is produced using the viral genome. This includes modifications to polypeptides encoded for by the virus.

Components include polypeptides comprising the virus, such as envelope proteins, processing enzymes, structural proteins, nucleic acid synthesis enzymes, glycoproteins, carbohydrates, lipids, antigens or antigenic fragments of the virus, etc. Also included are nucleic acid fragments of the whole genome, including fragments comprising complete gene sequences, control sequences, etc.

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Components includes one or more of the over about 198 open reading frames (ORF) and about 268 genes that have been identified in vaccinia and other poxvirus. Components include one or more of the genes and products thereof described in, but not limited to, Antoine et al., Virology, 244:365-396, 1998, and Goebel et al., Virology, 179(1):247-266, 1990 for vaccinia virus; Willer et al., Virology, 264(2):319-43, 1999 for Leporipoxvirus Shope fibroma virus (SFV); Cameron et al., Virology, 264(2):298-318, 1999 for myxoma virus; Shchelkunov et al., Virology, 297(2):172-94, 2002 for monkeypox virus; Shchelkunov and Totmenin, Virus Genes, 9(3):231-45, 1995 for variola, Massung et al., Virology, 201(2):215-40, 1994. For example, the polypeptide coding for the 17K myristylprotein, and which has amino acid sequence homology to gp120, can be used alone or in combination with other antigens, etc., in accordance with the present invention. See, e.g., Antoine et al., 1998; Barrett et al., Seminars in Immunol., 13:73-84, 2001. See, also Tables 1 (from Goebel et al., Virol., 179:247-266, 1990) and 2 (from Antoine et al., Virol., 244:365-396, 1998). Moreover, one or more of the aforementioned genes and open reading frames can be deleted from a vaccinia virus, e.g., to eliminate a toxic or other undesirable effect of an administered virus.

A useful composition can comprise one of the components of a poxvirus, including one or more of the components described in Tables 1 and 2. These can be individual purified and then combined into a therapeutic or prophylactic composition, or extracts can be prepared from viral particles and treated as desired. The individual components can be purified from the viral particles, or produced recombinantly, e.g., where a target gene is cloned, expressed in a host cell under conditions where the polypeptide is manufactured by the cell, and separating and purifying the polypeptide

accordingly to conventional methods. Components can also be administered as naked DNA. See, e.g., U.S. No. 6,413,942.

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The therapeutic and/or prophylactic effect achieved with the poxvirus can be independent of an immunological response to it. For example, the purpose of ordinary smallpox vaccination is to elicit an immune response by the host. This response is both humoral and cellular, involving the generation of specific antibodies and immune cells (such as T-cells, cytolytic or cytotoxic T lymphocytes, etc.) which protect a host from future invasion by the smallpox virus. While the present invention is not bound by any mechanism through which the poxvirus achieves its therapeutic and/or prophylactic effect, it can be mediated through a pathway separate from the immune response and not require cellular or humoral immunity. For example, poxvirus, or a component thereof, can directly block or inhibit the ability of a HIV to infect a cell. In this respect, the poxvirus, or component of it, acts as an antagonist, blocker, etc., of HIV's ability to infect target cells. HIV usually activates a G-proteincoupled signal pathway cascade. Poxvirus can interfere with this pathway or modify it such a way that the cell is more difficult to infect, thereby increasing its resistance to the HIV virus. Consequently, the effective amounts of a poxvirus, or component thereof, can differ from the amounts that are ordinarily used when the objective is to achieve a humoral and/or cellular immune response.

Vaccination with vaccinia can be associated with adverse reactions. Those at highest risk include, e.g., pregnant women, immunocompromised patients (e.g. HIV-positive), and persons who have atopic dermatitis or eczema. Strains which are attenuated or otherwise modified to reduce adverse effects are especially useful in accordance with the present invention, e.g., for administration to persons at risk for adverse effects.

Modified strains of vaccinia can be utilized that are deficient, mutated, engineered, etc., in one or more of the about 198 open reading frames (ORF) and/or about 268 genes that comprise vaccinia (depending on the strain or variant). In addition, genes can be inserted into vaccinia, including, one or more copies of a vaccinia gene of interest (e.g., 17K myristylprotein, vCCI), and/or genes coding for all or part of an HIV proteins, such as gp120 or gp40.

The present invention also provides methods of treating and/or preventing HIV infection in a subject in need thereof, comprising, e.g., administering multiple doses of a poxvirus, or components thereof, to a subject, wherein each dose is administered at a time interval from the previous dose, and are effective to maintain a therapeutic effect, or to maintain protection against HIV infection. As discussed above, a dose of the poxvirus, or component thereof, is the amount of virus which is useful for accomplishing the therapeutic or prophylactic effect. More than one dose can be administered to the subject in order to maintain the therapeutic efficacy of the treatment, or to maintain protection against HIV infection. For example, smallpox immunization is usually achieved by a single vaccination with a booster every 5-10 years. To maintain protection against HTV, more frequent vaccination can be used, e.g., multiple times a year, at least twice a year, yearly, every two years, every three years, more than once every less than five years, more than once every less than ten years, etc. The periods between the separate and sequential vaccinations can be referred to as "time intervals." These intervals can be spaced apart by any desired time period which is effective to maintain protection or therapeutic efficacy in treating an infected subject. The intervals can be predetermined or preset, where they are already specified, or they can be determined by monitoring the progress of a subject, e.g., using blood serum to measure poxvirus antibody titer, or HIV titer in an infected subject. The frequency of vaccination utilized to achieve efficacy may vary depending upon multiple factors, including, e.g., person-to-person variations in the immune system, the stage of HIV infection, the potency of the virus or vaccine, etc. and may be as often as every 3 months to once every 5 years.

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The present invention also provides methods of treating and/or preventing lentivirus infection in a subject in need thereof, comprising: administering an effective amount of a poxvirus or component thereof, wherein said amount is effective to treat and/or prevent lentiviral infection, with the proviso that a lentivirus nucleic acid, such as HIV, is not contained in the poxvirus genome. This excludes, e.g., a poxvirus which is utilized as a vector to administer HIV nucleic acid, such as when HIV nucleic acid is inserted into the poxvirus genome.

The present invention also provides methods of identifying a component of a poxvirus, or a poxvirus-associated agent, which interferes with HIV infection, and

components and agents identified thereby. Interfering with HIV infection indicates that the agent or component decreases, reduces, diminishes, lessens, etc., the ability of a susceptible cell or organism to become infected with HIV virus as compared to the same cell or organism in the same conditions, but in the absence of the agent or component. Interference with HIV infection can occur at any level, e.g., by blocking the ability of HIV to attach to its receptor(s) on a cell, by blocking the ability of HIV to be taken into a cell, by blocking viral function once inside the cell, by blocking viral infection, etc. The invention is not limited by the mechanism through which HIV interference is achieved. By interfering with HIV infection, the cell's or organism's resistance to HIV is increased.

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These methods can involve one of more of the following steps in any effective order, e.g., (1) contacting a cell or organism which is susceptible to HIV infection with poxvirus, or a component thereof, or a poxvirus-associated agent, (2) contacting said cell or organism with HIV under conditions effective for said HIV to infect said cell or organism, and, (3) (a) determining whether said cell or organism is resistant to HIV infection, whereby said agent is identified as interfering with HIV infection, or (3) (b) identifying the poxvirus, or component thereof, which confers resistance to HIV infection. The term "organism" as used herein indicates a fully-gestated animal.

The method can also involve a step of identifying the poxvirus, or a component thereof, as the agent which confers resistance to HIV infection. Identifying the poxvirus, or component thereof, which confers resistance to HIV infection, indicates that the poxvirus is positively determined or ascertained to provide protection or resistance against HIV. This indicates a positive result in the method.

Agents can be tested for their ability to interfere with HIV infection in any suitable system, including whole animals and cell culture. Animal cells useful in the present invention are those which are susceptible to HIV infection, i.e., they are capable of being infected by the HIV virus. They can be naturally-susceptible, or genetically-engineered to confer susceptibility, e.g., by expressing HIV receptor (CCR5, CD4, etc.), or by grafting on the human immune system. Any methods for testing whether a cell or organism is infected with HIV can be used, e.g., measuring

anti-HIV antibody titer (e.g., gp120 antibodies), reverse transcriptase protein or nucleic acid, or any other polypeptide or nucleic acid.

Any suitable animal model for testing the efficacy and dosage of a poxvirus (or component thereof) can be used in accordance with the present invention. These include, but are not limited to, e.g., SCID mice reconstituted with human immune 5 system components (e.g., peripheral blood lymphocytes) [e.g., Zhang et al., Proc. Natl. Acad. Sci., 93:14720-14725, 1996, using SCIC.bg mice], chimpanzees infected with HIV-1, macaque monkeys infected with SIV, HIV2, or chimeric SIV/HIV [e.g., Johnson, Curr. Opin. Immunol., 8(4):554-560, 1996], cats infected with feline immunodeficiency virus, HIV-1 transgenic mouse model [e.g., mice which have 10 integrated molecular clone pNL4-3 containing 7.4 kb of the HIV-1 proviral genome deleted in the gag and pol genes (Dickie et al., Virology, 185:109-119, 1991; transgenic mice carrying an HIV provirus, optionally with deletion of one or more HIV genes (Tinkle et al., J. Clin. Invest., 100(1):32-9, 1997)], HIV-1 transgenic rat model, human CD4 transgenic rat model, horse infected with EIAV, sheep infected 15 with visna virus, goats infected with CAEV, etc. See, also, The Retroviridae, J. A. Levy, ed., Plenum Press, 1993, e.g., Chapters 3, 4, and 5.

A vaccinia virus-associated agent is any substance which is produced in response to a vaccinia infection, or in response to inhalation, injection, ingestion, etc., of any vaccinia virus, or component thereof. This substance can be present in a culture medium in which cells exposed to vaccinia have been cultured, or can be present in blood serum when harvested from an organism exposed to vaccinia. The present invention provides compositions which comprise such substances.

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The invention also provides combinations of pharmaceutical agents for treating and/or preventing HIV, e.g., poxvirus, or a component thereof, and an agent which is used to treat HIV, such as a protease inhibitor or a reverse transcriptase inhibitor. Examples of the latter classes of drug, include, but are not limited to, saquinavir, ritonavir, indinavir, nelfinavir, amprenavir, lopinavir, atazanavir, fosamprenavir, tipranavir, AZT, ddI, ddC, ddT, 3TC, nevirapine, delavirdine, etc. The active agents can be present in the same dosage unit (e.g., a composition), or can be used as separate dosage units.

In addition, a poxvirus, such as vaccinia, can be administered in combination with HIV nucleic acid. The HIV nucleic acid can be physically joined to the poxvirus genome, or it can be administered as a separate component. For example, HIV nucleic acid (e.g., coding for gp120 or another viral antigen) can be administered at the same time as a poxvirus, but as a physically separated entity, or it can be administered at subsequent times after receiving only poxvirus) as part of a regimen for treating and/or preventing HIV infection.

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The present invention also provides methods of making a poxvirus composition for conferring resistance to HIV infection or treating HIV infection,, comprising, one or more of the following steps in any effective order, e.g., preparing a composition comprising poxvirus, or a poxvirus component thereof, and/or identifying that the poxvirus, or component thereof, confers resistance to, or treats, HIV infection. As mentioned earlier, the identifying step indicates obtaining a positive result in finding that the poxvirus (e.g., vaccinia), or component thereof, provides resistance, protection, treatment, etc., against the HIV virus.

The preparation of a poxvirus composition can be carried out routinely, e.g., according to conventional methods used for vaccine manufacture. Preparing includes culturing poxvirus, isolating poxvirus, putting poxvirus into a form suitable for administration (oral, injection, nasal, etc.), making poxvirus components recombinantly, etc. The prepared poxvirus (or components of it) can be assayed for its ability to confer resistance to HIV infection to an organism challenged with it or provide a therapeutic effect. By this, it is meant that a sample of the prepared composition is tested to determine its titer, concentration, potency, etc., in making a subject, to whom it is administered, "resistant" to the HIV virus, or for its therapeutic effect. The assay step can be carried out on every batch, or only selected batches, etc. A purpose of this step is, e.g., to confirm that the manufactured poxvirus possesses an anti-HIV activity for which it is to be administered. Any suitable assay or testing method can be utilized, e.g., in vitro methods of evaluating its efficacy or potency. For instance, the determining step can involve, e.g., challenging said organism, or cells derived from it, with infectious HIV, and detecting the expression in saidorganism or cells of gp120, HIV reverse transcriptase, p24, infectious HIV particles, and/or HIV nucleic acid. By "challenge" it is meant the cells or organism are placed

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in contact wit the HIV virus under conditions which are effective to become infected by it. These conditions will vary, depending upon how the assay is specifically accomplished.

When poxvirus is administered to a host, it can elicit a cellular response that is responsible or associated with the host's subsequent ability to resist HIV infection and/or treat HIV infection. This response can be measured, and used as index or marker to assess the efficacy of the poxvirus, and/or to determine effective amounts of it for the desired purpose (i.e., treating or preventing HIV infection). The appearance of one or more of the following "markers" can be modulated (e.g., elicited, stimulated, down-regulated, up-regulated, etc) by poxvirus, and associated with its anti-HIV effect, thereby making the marker useful as an indicator of poxvirus efficacy. By the term "marker," it is meant any measurable response to a poxvirus, including its effect on HIV's ability to infect and replicate in a cell, as well as on the host's immune system and the cells which comprise it. These markers, include, but are not limited to, one or more of the following agents, activities, responses, pathways, etc.:

- CD4 expression, e.g., measuring the amount of CD4 present in a cell-type that is susceptible to HIV infection
- HIV coreceptor expression, e.g., CCR5 or CXCR4 chemokine receptor, including its cell-surface expression
  - Cytokine receptors
  - Virus-specific CTLs (cytolytic or cytotoxic T-cells, including CD8+ T-cells) which are capable of lysing HIV infected cells (cells can be co-infected with poxvirus and HIV, or infected by HIV alone)
- 25 CD8 cells
  - Cytokines, including mediators and regulators of innate immunity, such as interferons, type I interferon, interleukins, interleukin-15, interleukin-12, tumor necrosis factor, interleukin-1, interleukin-6, interleukin-10, etc.; and mediators and regulators of specific immunity, such as interleukins, interleukin-2, interleukin-4, transforming growth factor-beta, interferon-gamma, lymphotoxin, interleukin-5, etc.
  - Chemokines (a large family of structurally homologous cytokines, that, e.g., stimulate leukocyte motility and directed movement), including, but not limited to,

the C-C and C-X-C families. Examples of chemokines, include, but are not limited to, e.g., interleukin 8, Gro, platelet basic protein, epithelial-derived neutrophil attractant 78, platelet factor 4, interferon-gamma-induced protein 10, stromal cellderived factor-1, monocyte chemotactic proteins 1, 2, and/or 3, RANTES, monocyte inflammatory protein 1-alpha and 1-beta ("MIP"), eotaxin, lymphotaxin, etc.

- Th1/Th2 phenotype and cytokine secretion pattern. Effector T-cells (e.g., CD4+ helper T-cells) can be categorized, on the basis of the cytokines they secrete, into Th1 and Th2 cells. Th1 cells secrete, e.g., interferon-gamma, lymphotoxin-alpha, TNF-beta, IL-2, IL-10, and CCR5 ligands, such as RANTES and MIPS. Th2 cells secrete, e.g., IL-4, IL-5, IL-6, IL-9, IL-10, IL-13, etc. Th1 and Th2 cells also include 10 resting, but polarized T-cells (i.e., committed to a Th type). In addition to cytokine production profiles, there are a number of cell surface markers that can be used to differentiate between Th1 and Th2 subtypes. For example, Th1 cells express both components of IL-12 receptor chains (beta1 and beta2), while Th2 cells exhibit IL-12R-beta1. Th2 cells exhibit both IFN-gamma receptor chains (a and b), while Th1 15 cells express IFN-gamma-R-alpha. Th2 cells appear to express a fully functional IL-1 receptor, and ST2L/T1, an IL-1R-like molecule, is found on Th2 cells. Chemokine receptors CXCR-3 and CCR-5 are also characteristic of Th1 cells, while CXCR-4, CCR-3, CCR-4, CCR-7 and CCR-8 are associated with Th2 cells. CD30, a member of the TNF superfamily, is associated with Th2 cells. The Th1/Th2 pattern can be polarized by poxvirus administration, resulting in a phenotype that favors the secretion, etc., of cytokines that inhibit HIV infection and/or render cells resistant to infection. One or more of the aforementioned molecules can be utilized as markers of poxvirus efficacy

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- Antibodies that specifically recognize HIV, e.g., neutralizing antibodies
- Antibodies that specifically recognize poxvirus
- Complement control protein. Vaccinia virus encodes a secreted complement control protein (VCP, 35-kDa) protein with sequence homology to the SCRcontaining complement control protein superfamily. It binds C3b and C4b, and interferes with the complement cascade by providing cofactor activity for the cleavage of C3 and C4 by factor I, and by accelerating the decay of the C3 converse of both the alternative and, more effectively, the classical pathway of complement

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activation. VCP may suppress the complement system or their receptor expression, rendering the host less susceptible to the complement-enhancement of HIV infection

- Activation state of a cytokine receptor, e.g., CCR5 receptor or other HIV chemokine coreceptor. For example, poxvirus can interfere with CCR5 activation after HIV binding, e.g., by modulating tyrosine kinase feedback pathways
- One or more of the vaccinia proteins listed in Tables 1 and 2. This includes any poxvirus-encoded protein that specifically interferes with CCR5/CD4/gp120 interactions, including, e.g., vaccinia encoded CC chemokine binding proteins and/or IFN-gamma receptor-like protein
  - RNA interference with HIV expression/replication in infected cell
  - Alpha-defensins 1, 2, and/or 3
- Soluble factors including those produced by CD8+ lymphocytes and sometimes referred to as CAF
- Interference with the HIV life cycle, including viral entry, import into the host cell nucleus, viral integration into host genome, Rev-dependent and Rev-independent transport from the host nucleus, replication, gene expression, RNA splicing, etc
- Inhibiting HIV replication, including its ability to make copies of itself in the cell, and for productive viral particles to be extruded into the blood
- Inhibiting the ability of HIV to infect a cell, e.g., to bind to CD4 and/or its coreceptor, for the envelope protein to fuse with the host cell membrane, etc.
- Modulating gene expression of the HTV virus, including modulating regulatory genes (e.g., tat and rev), accessory genes (e.g., vif, vpu, vpr, and nef), structural genes (e.g., gag, pol, and env), inner core polypeptides (e.g., gag, p17, p24, p7, and p9), viral enzymes (pol, reverse transcriptase, protease, and integrase), and envelope proteins (e.g., env, gp120, and gp41). The phrase "gene expression" is used broadly to mean any step in the pathway from viral RNA to protein synthesis, and therefore includes all regulatory processes, transcription, translation, polypeptide processing, etc.
- Modulating activity of a HIV encoded polypeptide, including, tat, rev, vif, vpu, vpr, nef, gag, p17, p24, p7, p9, pol, reverse transcriptase, protease, integrase, env, gp120, gp41, etc.

- Modulating viral regulatory sequences, such as RRE, cis-acting repressive sequences (CRS), and inhibitory/instability RNA sequences (INS)

- Any cell or tissue of the immune system, including, but not limited to, lymphocytes, B lymphocytes, T lymphocytes, helper T cells, cytotoxic (or cytolytic) T cells ("CTL), natural killer (NK) cells, naïve T cells, memory T cells, CD4+ helper T cells, CD8+ CTLs, monocytes, macrophages, antigen-presenting cells (APCs), dendritic cells, granulocytes, etc.

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The present invention also provides kits comprising a poxvirus. For example, a kit for preventing HIV infection, comprising: an effective amount of a poxvirus, and instructions for administering an effective amount of said poxvirus to a subject to prevent HIV infection; and a kit for treating HIV infection, comprising: an effective amount of a poxvirus, and instructions for administering an effective amount of said poxvirus to a subject to treat HIV infection. The instructions can provide any information that is useful for directing the administration of the poxvirus for the desired purpose.

The present invention also provides methods of advertising, licensing, selling, purchasing, etc., a poxvirus for the purpose of treating and/or preventing HIV infection. Methods can comprise, one or more of the following steps in any effective order: e.g., displaying information (a) comprising instructions for administering a poxvirus for treating and/or preventing HIV infection or (b) comprising a description of the use of poxvirus for treating and/or preventing HIV infection, in a printed or computer-readable medium (e.g., on the Web, Internet, personal computer, server, etc); offering for sale a poxvirus for treating and/or preventing HIV infection in a printed or computer-readable medium; accepting an offer to purchase poxvirus for said use in a printed or computer-readable medium.

### **EXAMPLES**

The following experiments were performed in the laboratory of Dr. Beda Brichacek and Dr. Michael Bukrinsky of the Department of Microbiology and Tropical Medicine, The George Washington University, Washington D.C. 20037.

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### Methods

## Subject selection and specimen collection.

Twenty subjects were chosen for inclusion in the study. Ten subjects had been immunized with vaccinia within the previous 3 to 6 months, and ten subjects had never been immunized with vaccinia. All subjects were healthy and had a negative HIV test within the previous year. No subjects of northern European descent were used in order to avoid the potentially complicating factor of including a subject who might be homozygous for the CCR5-delta32 mutation. Two tubes of heparinized blood and 1 serum separator tube were collected. All blood samples from all subjects were drawn within 6 hours of each other, and were immediately processed to separate the PBMCs using standard methods of Ficoll-Hypaque centrifugation.

### Cell culture preparation.

PBMCs were centrifuged at 1200 rpm for 11 minutes and resuspended in RPMI tissue culture medium + 10% fetal calf serum + 10  $\mu$ g/ml gentamicin at a concentration of about 1-3 x 10<sup>6</sup> cells/ml with a final concentration of 2 x 10<sup>6</sup> cells/culture. Cell cultures were incubated in a CO<sub>2</sub> incubator. On the second day, one of the utilized strains of HIV was mixed with either culture medium or serum from each individual subject and incubated on ice for 7 hours after which 175  $\mu$ l of each mixture was added to the autologous cell cultures. The next day 1 ml of cell culture media was added and the cultures were incubated for 5 hours to dilute the viral inoculum and to allow the virus to detach. The supernatant was carefully aspirated and 1 ml of fresh media was added before the cultures were spun down at 1000 rpm for 7 minutes. The supernatant was again aspirated and 2 ml of fresh media was added to each culture. 150  $\mu$ l of supernatant for RT analysis was aspirated from each culture tube on days 2, 5, 8 and 10, and if needed, up to an additional 1 ml was aspirated and replaced with fresh media. On day 2, PHA was added to the tubes of

culture series F to act as a cell activator. On day 5, 2 ml of supernatant was removed from each of tubes of culture series F and replaced with 2 ml media + human serum + IL-2.

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# 5 Reverse Transcriptase (RT) analysis.

The measurements of viral replication were performed by standard RT assays using tritiated thymidine as described in numerous articles in the scientific literature. See, e.g., Rey et al., *Virology*, 181(1), 165-71, 1991.

### 10 Results

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All results are based on RT analysis using tritiated thymidine, and are given in counts per minute (CPM).

Culture Series A, the control, demonstrated no viral replication in any cultures.

15 Culture Series B (without serum; Fig. 1A) demonstrated a significant reduction of viral replication in most cultures from vaccinated subjects when compared to unvaccinated subjects. Two subjects (1 and 10) showed a complete lack of viral replication, comparable to the controls in culture series A. One subject was excluded from all analyses when it was subsequently discovered that the subject had had a highly anomalous reaction to the vaccinia immunization with recurrent skin lesions for months afterward. This suggested an inadequate immune response to the vaccinia, and this subject correspondingly did not show any protection against HIV in cell culture, demonstrating viral replication comparable to unvaccinated subjects.

Culture Series C (with serum; Fig. 1B) also demonstrated a significant reduction of viral replication in most cultures from vaccinated subjects, when compared to unvaccinated subjects. The same two subjects (1 and 10) noted in culture series B also had no demonstrable viral replication, comparable to the controls in culture series A. The addition of autologous serum in culture series C further enhanced the difference between vaccinated and unvaccinated subjects when compared to culture series B (no serum).

Culture Series D, E and F, using the T-cell (CXCR4) tropic HIV (Fig. 1C), demonstrated no difference between vaccinated and unvaccinated subjects, including

the two subjects (1 and 10) who were resistant to infection by the macrophage (CCR5) tropic HIV in culture series B and C. As stated in the methods section, care was taken in the selection of subjects to avoid those of northern European descent who might be homozygous for the CCR5-delta32 mutation, so this cannot be an explanation for the described resistance. There was also no difference noted between the addition of serum and no serum (cultures D and E).

#### Discussion

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By at least day 10, there is a statistically significant difference between the vaccinated and non-vaccinated subjects in culture series B and C (p=.035 and .013 respectively) that increases by day 13 (p=.017 and .008 respectively), indicating a resistance to infection by HIV in the vaccinated subjects (Fig. 1). Subjects 1 and 10 demonstrated total resistance to macrophage (CCR5) tropic HIV infection in both culture series B and C, with RT measurements equal to the non-HIV infected control (culture series A). The fact that the same result was achieved in both sets of cultures, while infection was easily achieved with the T-cell (CXCR4) tropic HIV in cultures D, E and F, indicate these finding were not the result of laboratory error.

Without further elaboration, it is believed that one skilled in the art can, using the preceding description, utilize the present invention to its fullest extent. The following preferred specific embodiments are, therefore, to be construed as merely illustrative, and not limitative of the remainder of the disclosure in any way whatsoever. The entire disclosure of all applications, patents and publications, cited above and in the figures are hereby incorporated by reference in their entirety, including of U.S. Provisional Application Nos. 60/491,258 filed July 31, 2003, 60/493,767 filed August 11, 2003, 60/496,908 filed August 22, 2003, and 60/501,832 filed September 11, 2003.

TABLE 1
THE OPEN READING FRAMES OF VACCINIA VIRUS

_		<u>ation</u> [	<u>s</u>	ize		•
Gene <sup>a</sup>	Start	Stop <sup>b</sup>	aa	MC	Characteristics <sup>d</sup>	References
C23L*	5008	4277	244	26.4		
					Nonessential; B29R Acidic <sup>e</sup> (4.2)	Perkus, et al. (1990b)
C22L*	6113	5748	122	13.6	Nonessential; B28R Hydrophobic N-terminus	Perkus, et al. (1990b)
C21L*	6815	6477	113	13.4		· Portug of all (2000)
C20L*	7132	6824	103	12.5	Nonessential; B26R	Perkus, et al. (1990b) Perkus, et al. (1990b)
				*	Basic (9.0)	101x43/ et al. (1990b)
C19L*	7856	7080	259	30.5	Nonessential; B25R	Perkus, et al. (1990b)
C18L*	8693	8244	150	17.5	Hydrophobic N-terminus Nonessential; B24R	Perkus, et al. (1990b)
C17L*	9947	8790	206	44.0	Acidic (4.8)	, ,
C16L*	10539	9997		44.9	Nonessential; B23R	Perkus, et al. (1990b)
C15L*	11153			21.0	Nonessential; B22R	Perkus, et al. (1990b)
C14L	.12212	10881		10.5	Nonessential; B21R	Perkus, et al. (1990b)
CIAD	.12212	11967	82	9.3	Nonessential Basic (9.2)	Perkus, et al. (1990b)
C13L	12510	12316	65	7.4	Nonessential	Perima at al (1000)
C12L	1 2722	12675	252		Acidic (4.0)	Perkus, et al. (1990b)
CIZL	13733	12675	353	40.4	Serine Protease Inhibitor	Kotwal and Moss (1988b)
					Nonessential	Perkus, et al. (1990b)
Clir	14178	14603	142	15 0	Acidic (4.8)	
	14170	14003	142	15.8	Growth Factor	Blomquist, et al. (1984);
					·	Brown, et al. (1985); Reisner (1985)
					Nonessential	Buller, et al. (1988); Perkus
					EGF-like type A domain	et al. (1990b)
					Hydrophobic C-terminus	• •
ClOL	15754	14762	331	38.5	Nonessential	Dominion
					Acidic (4.5)	Perkus, et al. (1990b)
<b>29L</b>	18136	16235	634	74.7	Nonessential	Parloug of al (1000)
						Perkus, et al. (1990b);
CSL	18733	18182	184	21.6	Nonessential	Kotwal and Moss (1988b)
		•				Kotwal and Moss (1988b);
					Acidic (4.4)	Perkus, et al. (1990b)
27L	19257	18808	150	18.0	Nonessential	Kotwal and Moss (1988b);
-						Perkus, et al: (1990a,b)
					Host range function	Perkus, et al. (1990a)
26L	19939	19487	151	17.4	Nonessential	Kotwal and Moss (1988b);
					•	Perkus, et al. (1990b)
					Acidic (4.8)	101 Add, et al. (1990b)
SL	20680	20069	204	24.5	Nonessential	Kotwal and Moss (1988b);
					<del>-</del>	Perkus, et al. (1990b)
					Acidic (4.8))	Terkus, et ar. (1990b)
24L	21693	20746	316	37.2	Nonessential	Kotwal and Moss (1988b);
						Perkus, et al. (1990b)
:3L	22551	21763	- 263	28.6	Nonessential	Kotwal and Moss (1988a,b);
			-			Perkus, et al. (1990b)
	•				C4B binding protein	(25500)
					homolog; virokine	Kotwal and Moss (1988a)
2L .	24156	22621	512	59.2	Nonessential	Kotwal and Moss (1988b);
						Perkus, et al. (1990b)
			•		Hydrophobic N-terminus	
1L	24900	24229	224	26.4		Kotwal and Moss (1988b);
						Perkus, et al. (1990b)
					Basic (9.0)	

Reprinted from *Virology*, Vol. 179, S. J. Goebel, G. P. Johnson, M. E. Perkus, S. W. Davis, J. P. Winslow and E. Paoletti, "The Complete DNA Sequence of Vaccinia Virus", pgs. 247-266 (1990), with permission from Elsevier.

TABLE 1—Continued

	Transl		Si	ze _		
Gene <sup>a</sup>	Start	Stopb	aa	MC	Characteristics	References
						h
						•
I1L	25240	24890	117	14.0	Nonessential	Kotwal and Moss (1988b);
					Virokine	Perkus, et al. (1990b)
					Acidic (4.2)	Kotwal and Moss (1988a)
12L	25886	25362	175	20.8	Nonessential	Kotwal and Moss (1988a,b);
						Perkus, et al. (1990b)
11L	27346	25931	472	54.2	Nonessential	Perkus, et al. (1990b)
	00000	02202			Homology to K1L	Perkus, et al. (1990a)
12L	27986	27327	220	25.1	Nonessential Hydrophobic N-terminus	Perkus, et al. (1990b)
					nydrophobic w cerminds	
KIL	28975	28124	284	32.6	Host range function	Gillard, et al. (1986);
	•				Nonessential	Perkus, et al. (1989) Perkus, et al. (1990b)
					Monessential	rerkus, et al. (1990b)
K2L	30313	29207	369	42.3		Boursnell, et al. (1988)
					Nonessential Basic (9.3)	Perkus, et al. (1990b)
K3L	30629	30366	88	10.5	Nonessential	Perkus, et al. (1990b)
					Basic (9.3)	
(4L	31955	30684	424	48.9	Translation initiation fa Homology to F13L	Boursnell, et al. (1988)
				,,,,,	Nonessential	Perkus, et al. (1990b)
K5L	32497	32090	136	15.2		Perkus, et al. (1990b)
(6L	32764	32522	81	9.1	Basic (10.2) Nonessential	Perkus, et al. (1990b)
K7R	32903	33349		17.5	Nonessential	Perkus, et al. (1990b)
					Acidic (4.4)	
		*			Hydrophobic C-terminus	
F1L	34097	33420	226	26.4	Nonessential	Perkus, et al. (1990b)
					Acidic (4.4) Hydrophobic C-terminus	
F2L	34552	34112	147	16.3	Retroviral protease	Slabaugh and Roseman (1989
					Nonessential	Perkus, et al. (1990b)
F3L	36018	34579	480	55.7	dUTPase Nonessential	Perkus, et al. (1990b)
74L	36988	36032	319	37.0	Ribonucleotide reductase	Slabaugh, et al. (1988)
					(small subunit)	
					Nonessential Acidic (4.6)	Perkus, et al. (1990b)
75L	37985	37023	321	36.5	Multiply hydrophobic	
F6L	38239	38018	74	8.6	Acidic (4.1)	
77L 78L	38533 38878	38258 - 38684	92 65	7.8	- (Lys-Asn) <sub>9</sub> Basic (9.9)	
79L	39576	38941	212	23.8	Hydrophobic C-terminus	
FIOL	40882			52.2	Protein kinase 2nd signat	ure
F11L F12L	41969 43919	40908 42015		39.7 73.2	<del>-</del>	
713L	45079	43964		41.8	Envelope antigen	Hirt, et al. (1986)
F14L	45318.		73	8.3	Acidic (2.9)	•
F15L F16L	46068 46770	45595 46078		18.6 26.6	Basic (9.5) Basic (9.6)	
F17R	46833	47135		11.3	Basic (9.8)	
E1L E2L	48574 50784	47138 48574		55.6 85.9	<del>-</del> <del>-</del>	
22L 23L	51483	50 <del>9</del> 14		21.5	Acidic (4.9)	•
E4L	52318	51542		29.8	Acidic (4.9)	
					Transcription factor	

-22-

TABLE 1—Continued

Gene <sup>a</sup>	<u>Transl</u> Start	ation Stop <sup>b</sup>		M <sub>r</sub>	Characteristics	References
<u> </u>						
ESR	, 52395	53387	331	30.1	(ts: C19??) <sup>f</sup>	Condit, et al. (1983)
E6R	53527	55227		39.1 66.7	Basic (9.8)	
E7R	55314	55811		19.5	-	
E8Ř	55939	56757		31.9	Basic (9.3)	
E9L	59787	56770		117.0	DNA Polymerase	Earl, et al., 1986
					ts: C42, NG26;	2227, 66 81., 1986
					PAAr, Aphidicolin	Traktman, et al. (1989b)
E10R	59819	60103	95	10.8	DNA polymerase family B s	ignature
E11L	60490				_	
		,				
OlL	62477	60480		77.6	Leucine Zipper Motif	
O2L	62851	62528	108	12.4	Glutaredoxin	
IIL .	62025	63000	2.10	25 0		•
IZL	63935 64163	63000 63945		35.8 8.4	The decay be below as the second	·
	04103	00743	/3		Hydrophobic C-terminus Acidic (3.9)	
I3L	64973	64167	269	30.0	- (3.9)	
I4L	67371	65059		87.0	Ribonucleotide reductase	Schmitt and Stunnenberg (1988)
					(large subunit)	Tengelsen, et al. (1988)
			•		Nonessential	Perkus, et al. (unpublished)
						Child, et al., (1990)
	,				Divalent Fe-S ferredoxin	
ISL	67637	67401	79	8.7	binding region signature	
I6L	68804	67659		43.4	Basic (9.9) Basic (9.2)	
I7L	70068	68800	423	49.0	-	
IBR .	70074	72101	676	77.6	ATP/GTP binding motif A	
G1L	71003	20111			-	
G2R	73883 74209	72111 74868	591	67.9	<b>-</b>	
G3L	74215	73883		25.7 12.8	The decomposition of the same of	
G4L	75215	74844	124	14.0	Hydrophobic N-terminus Acidic (4.8)	
G5R	75218	76519		49.9	Acidic (4.8)	
G6R	76723	77217	165		_	·
G7L	78300	77188	371	41.9	-	
G8R	78331	79110	260	29.9	<del>-</del>	
G9R	79133	80152	340	38.8	Hydrophobic C-terminus	
LlR	80156	80905	250	27.3	Hardwannah da a a a a a a a a	
L2R	80940	81200	87	10.2	Hydrophobic near C-termin	us
L3L	82245	81196		40.6	Multiply hydrophobic	
L4R	82270	83022	251		Structural protein, VP8	Yang, et al. (1988)
L5R	83035	83418	128	14.0	Basic (10.0)	11.37 00 41. (1300)
מוד.	02270	02026				
JlR J2R	83378 83855	83836 84385		17.8	mbaar 1 a 1 a 1 a 1	•
		04303	1//	20.1	Thymidine kinase	Weir and Moss (1983); Hruby et
					Nonessential	al. (1983)
					ATP/GTP binding motif A	Mackett, et al. (1982)
					/GIT DINGING MOCIL A	
J3R	84454	85452		15.2	Basic (10.0)	
J4R	85370	85924	185	21.3	RNA Polymerase subunit	Broyles and Moss (1986)
					ts: C7, C20	Hooda-Dhingra, et al. (1989);
757	95400	06000				Thompson, et al. (1989)
J5L J6R	86403 86510	86005		15.2	Hydrophobic C-terminus	
- OIK	20310	90367	1286	146.8	RNA Polymerase subunit	Broyles and Moss (1986)
					ts: E8, E13, E72	Ensinger (1987)
					C51, C53, C65	Hooda-Dhingra, et al., (1989); Thompson, et

TABLE 1—Continued

Gene <sup>a</sup>	<u>Transl</u> Start	ation Stopb	Si	ze M c	Characteristics	Poforongo
				M <sub>C</sub>	OWNT OCCEL TRETTER	References
H1L	90882	90370	171	19.7	Basic (9.6)	
H2R	90896			21.5	Hydrophobic N-terminus	
H3L	92442	91471	324		Multiply hydrophobic	
H4L	94830			93.6	- warerbil widtobuonic	
H5R	95016	95624		22.3	-	
H6R	95628	96569		36.7	Basic (10.0)	
H7R	96609				DNA topoisomerase	Shuman and Moss (1987)
			146	16.9	_	•
DIR	97093	99624	844	96.7	mRNA capping enzyme (small subunit)	Morgan, et al. (1984)
D2L	100026	99589	146	16.9	ts: E52, E94	Seto, et al. (1987)
D3R	100019	100729		28.0	ts: C5, C35	
D4R		101385		25.0	-	Seto, et al. (1987)
D5R		103774		90.0	ts: C17, C24, E69	Seto, et al. (1987)
D6R	103818	105728	637	73.8	ATP/GTP binding motif A Early transcription	Broyles and Fesler (1990)
					factor subunit ts: C46, E93	Seto, et al. (1987)
					Hydrophobic N-terminus	• •
D7R	105758	106240	161	17.9	RNA polymerase subunit	Ahn, et al. (1990)
					ts: C21, E45	Seto, et al. (1987)
					Acidic (4.5)	
DSL	107120.	106209	304	35.3	Carbonic anhydrase	Niles, et al. (1986)
					Transmembrane	Niles and Seto (1988)
					Cell surface binding	Maa, et al (1990)
					Multiply hydrophobic	
					Basic (9.1)	•
D9R		107800		25.0	<del>-</del>	•
DIOR		108543		28.9	-	
DllL	110442	108550	631	72.4	NTPase	Rodriguez, et al. (1986);
						Broyles and Moss (1987)
					ts: C36, C50, E17	Seto, et al. (1987)
D12L	111740	110400	207		Basic (9.0)	
DIZL	111340	110480	287	33.4	mRNA capping enzyme (small subunit)	Niles, et al. (1989)
DI3L	113026	111374	551	61.9	ts: C33, C43, E101	Seto et 27 (1997)
					Rifampicin resistance	Seto, et al. (1987) Tartaglia and Paoletti (1985);
						Baldick and Moss (1987)
					Acidic (5.0)	baldick and Hoss (1987)
AlL'	113502	113053	150	17.0		
A2L		113526		26.3	_	
A3L		114441		72.6	Major core protein P4b	Rosel and Moss (1985)
A4L		116428		30.8	Acidic (4.6)	Moser and Moss (1905)
A5R		117799		19.0	Acidic (4.2)	
A6L		117802		43.1		
A7L		118944		82.3	Early transcription	Gershon and Moss (1990)
A8R	121127	121990	200	33.6	factor subunit	
A9L		121989			_	
A10L		121989		11.1 102.3	-	Man Main and Mikkely (1000)
AllR		125929		36.1	Major core protein P4a Hydrophobic C-terminus	Van Meir and Wittek (1988)
Al2L	126512	125937	192	20.5	Acidic (4.7) Basic (10.1)	
A13L		126539	70	7.7	Basic (9.7)	
A14L		126859		10.0	- (3.7)	
A15L		127299		11.0		
A16L		127567		43.6	Hydrophobic C-terminus	
A17L		128706	203		Hydrophobic center	
· <b>-</b>				22.0	Acidic (4.1)	
A18R	129329	130807	493	56.7	Basic (9.3)	
				,		

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TABLE 1-Continued

	Trans	lation	S	ize		
Gene <sup>a</sup>	Start		aa	M <sub>C</sub>	Characteristics	References
	<del></del>	<del></del> .		<u> </u>		vererences
•						
Al9L	131024	130794	77	8.3	_	
A20R		132654			_	
A21L	131378	131028			Hydrophahia V	•
A22R		133147		-	Hydrophobic N-terminus	
A23R		134315		44.6	Basic (9.9)	
A24R		137806		137 /	Days	
		10.000	7704	133.4	RNA polymerase subunit; ts: C27, C29, C32,	Hooda-Dhingra, et al. (1990)
A25L	120011	122012	~-		C47, C62 Leucine Zipper Pattern	Hooda-Dhingra, et al, (1990)
12 JL	130011	137817	65	7.5	A-type inclusion protein (cowpox virus)	Funahashi, et al. (1988);
A26L	138948	137983	322	37.3	Acidic (3.3) A-type Inclusion protein	Funchashi at al case.
			•	•	(cowpox virus) Basic (9.2)	Funahashi, et al. (1988);
A27L	139330	139001	110	12.6	Fusion protein	Rodel
A28L	139771	139334		16.3		Rodriguez & Esteban (1987)
<b>A29L</b>	140689	139775		35.4	-	•
A3OL	140885	140655	77		Basic (9.9)	
A31R		141416		14.2	Basic (9.0)	
					Pibonual comments in Day 1	
A32L	142288	141389	300	34.4	Ribonucleoprotein RNA-bin Basic (9.2)	ding region signature
A33R	142316	142870	185	20.5	ATP/GTP Binding motif A	
A34R		143400		19.5	Pagin (10 1)	•
A35R	143447	143974		20.0	Basic (10.1)	•
A36R ´	144044	144706		25.1	Acidic (4.0)	
A37R		145561		29.9	Acidic (4.4)	
138L		145848				
139R		147903		31.6	Multiply hydrophobic	
140R		148435		45.7	- <del>-</del>	
141L		148499		19.3	Hydrophobic N-terminus	
142R		149732		25.1	Acidic (4.8)	
1721	143334	149/32	133	15.0	Basic (9.9) <i>Profilia</i>	
143R	149773	150354	194	22.6	- ,	
44L	151733	150696		39.4	3B-Hydrovy-5-one storoid	dah
145R		152154		13.8	3ß-Hydroxy-5-ene steroid Superoxide dismutase	aenyarogenase
46R		152788		24.7	- deroxide dismucase	t
47L		152959		28.3	Basic (10.0)	
48R		154400		23.2	Thymidylate kinase	
÷					ATP/GTP binding motif A Acidic (5.0)	Smith, et al. (1989a)
49R	154451	154936	162	18.8	Acidic (3.9)	
SOR	154972			63.4	DNA Ticaco	
			332	UJ,4	DNA Ligase	Colinas, et al. (1990); Smith, et al. (1989a); Kerr
					Nanagaah : - 1	and Smith (1989)
51R	156683	157694	324	37.7	Nonessential	Colinas, et al. (1990)
52R	157757	158324	100	22.7	Nonessential	Davis, et al. (unpublished)
.53R	158635				Hydrophilic N-terminus	
.54L				12.0	Nonessential	Davis, et al. (unpublished)
41.	158743	158474	90	10.8	Basic (10.4)	
cen	750445				Nonessential	Davis, et al. (unpublished)
.55R	159442	161133		64.7	Nonessential	Davis, et al. (unpublished)
56R	161186	162130	315	34.8	Nonessential	Shida, et al. (1987)
					Hemagglutinin Hydrophobic C-terminus	Shida (1986)
					3-111-13-5	
57R	163000	162730			Acidic (3.9)	

TABLE 1-Continued

а		ation b		.ze		
Gene <sup>a</sup>	Start	Stop <sup>b</sup>	aa	W.c	Characteristics	References
31R	162884	163783	300	34.3	ts: C2, C3, C25	Traktman, et al. (1989a)
					Protein Kinase	Howard and Smith (1989)
				•	Basic (9.1)	(4000)
32R	163876	164532	219	24.6	_	
33R	164571	164942		14.4	Acidic (4.7)	
34R		167276		65.3	-	
35R		168333		35.1	Multiply hydrophobic	
,,,,		10000	J.,	JJ.1	Acidic (4.4)	
					Complement control protein	
						ns
- < 5	160477	100000		20.1	C3L homologue	
36R		168950	173			
37R		169536	182		Hydrophobic N-terminus	
38R		170409		31.2	Hydrophobic N-terminus	
39R		170729	77	8.8	<del>-</del>	
310R		171192		18.9	<u>-</u>	
311R	171267	171530	88	9.9	Acidic (3.6)	•
			_		M(DT) <sub>Q</sub> DVTNV	
B12R		172448		33.4	Protein Kinase	Howard and Smith (1989)
313R	172562	172909	116	12.8	Hemorrhage-inducing	Pickup, et al. (1986)
					Serine Protease Inhibitor	Kotwal and Moss (1989);
			•		Nonessential	Perkus, et al. (1990b)
					Acidic (4.6)	•
B14R	172887	173552	222	24.9	Hemorrhage-inducing	Pickup, et al. (1986)
					Serine Protease Inhibitor	Kotwal and Moss (1989)
					Nonessential	Perkus, et al. (1990b)
					Acidic (4.3)	101AUD, CC 41. (13302)
815R	173632	174078	149	17.4	Nonessential	Perkus, et al. (1990b)
	1,0001	271070		47.4	Acidic (4.5)	101 Kdb, 01 d1. (1330b)
B16R	174272	175141	290	32.5	Nonessential	Perkus, et al. (1990b)
01011	1,42,1	1,2141	230	32.3	Kinase-related	reinus, et al. (1990b)
					transforming protein	•
B17L	176212	175193	340.	39.5	Nonessential	Perkus, et al. (1990b)
B17E B18R		178070		68.1	Nonessential	Perkus, et al. (1990b)
B19R		179203	353			Lervas' er at. (TAAAD)
アンド	1/0142	1/7/03	223	40.9	Hydrophobic N-terminus	Dawley et al. (1000)
205	170200	170000	100	1	Nonessential	Perkus, et al. (1990b)
320R	1/3700	179680	127	15.5	Nonessential	Perkus, et al. (1990b)
22154	100565	100055		30 5	Acidic (4.1)	N-13-1-1 (1000)
B21R*		180857		10.5	Nonessential; C15L	Perkus, et al. (1990b)
B22R*		181741		21.0	Nonessential; C16L	Perkus, et al. (1990b)
323R*		182948		44.9	Nonessential; C17L	Perkus, et al. (1990b)
324R*	183045	183494	150	17.5	Nonessential; C18L	Perkus, et al. (1990b)
					Acidic (4.8)	•
325R*	183882	184658	259	30.5	Hydrophobic N-terminus	
					Nonessential; C19L	Perkus, et al. (1990b)
326R*	184606	184914	103	12.5	Nonessential; C20L	Perkus, et al. (1990b)
					Basic (9.0)	•
327R*	184923	185261	113	13.4	Nonessential; C21L	Perkus, et al. (1990b)
328R*		185990		13.6	Nonessential; C22L	Perkus, et al. (1990b)
					Hydrophobic N-terminus	
B29R*	186730	187461	244	26.4	Nonessential; C23L	Perkus, et al. (1990b)
		TO / 40T		40.7	HOHEOBEILLIGI: CAJD	

<sup>&</sup>lt;sup>a</sup> Open reading frames enumerated as described in text.

<sup>\*</sup> Translation stop does not incude the three bases of termination codon.

<sup>&</sup>lt;sup>c</sup> M<sub>r</sub> values calculated for the nascent, unprocessed polypeptide chain are presented as kDa.
<sup>d</sup> Functions or activities indicated in bold type are known functions of vaccinia virus. Those indicated in *italics* have been identified in this study on the basis of similarity to existing proteins. All others are possible functions previously described by other investigators.

Acidic proteins: p/ < 5.0; basic proteins: p/ > 9.0. p/ presented within parentheses.

<sup>&</sup>lt;sup>7</sup>Temperature-sensitive mutants indicated by rs. Those first isolated by Condit et al. (1983) are prefaced with C; 1 begin with E. Mutant C19, while not localized to a particular open reading frame, appears to map in the vincinity of I

<sup>\*</sup> Open reading frames repeated in both left and right termini of genome.

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TABLE 2
Features and Homologies of Open Reading Frames of the Vaccinia MVA Strain

ORF <sup>1</sup>	STAD:			and Homologies of Open Re				
	STAR			name / (putative) function / homologies <sup>p</sup>	BLAST <sup>4</sup> expect		" HSS (%)	references
left t	6822	regi 136		201				
193R*	6412	130	14.9	35k major secr. protein chemokine receptor (f)				(Patel et al., 1990)
C23L		244		VAC (C23L/B29R)	6.0e-57	41/42	97	(Graham et al., 1997)
		253		VAR-P G3R	8.9e-51	46/49	93	(Goebel et al., 1990)
		246		CPX ORF B	5.6e-49	40/42	95	(Shchelkunov et al., 1995) (Hu et al., 1994)
		258		SFV T1 protein	2.5e-20	23/42	54	(Upton et al., 1987)
		260		Myxoma virus T1/35kDa	1.5c-14	. 21/42	50	(Graham et al., 1997)
002L/	7784	176	19.7	secr. TNF receptor (f)				
192Rb	7254	355		CPX errnB	5.le-71	76/83	91	(Upton et al., 1991a)
		348		VAR-BSH G2R	1.0e-66	73/83	87	(Hu et al., 1994) (Shchelkunov et al., 1995)
		326		Myxoma virus T2	4.9e-30	21/37	56	(Upton et al., 1991a)
		325 202		Rabbit fibroina Virus T2 CPX C4L	1.8e-28	17/36	47	(Upton et al., 1987)
		346		'HS TNF receptor protein	8.7¢-15	30/51	58	(Safronov et al., 1996)
CI9L		259		VAC (C19L/B25R)	1.9e-08 0.00026	14/26 16/19	53 84	(Heiler et al., 1990)
		277		human CD40L receptor	0.0015	11/24	45	(Goebel et al., 1990) (Stamencovic et al., 1989)
				30 matches to TNF receptors	< 0.39			(Stameneovic & at., 1989)
				and surface proteins				
003L/	3780	102	12.1	461				
191Rh	8472	102	1 4. 1	45k ank <sup>t</sup> -like protein (f1)				(Goebel et al., 1990)
CITL	1	386		VAC CI7L/B23R	1.3e-39	62/63		(0.11
004L/	9558	233	26.9	45k ank-like protein	1.36-33	02/03	70	(Goebel et al., 1990)
190Rb	8857			(f2)				(Goebel et al., 1990)
CI7L	1	386		VAC (C17L/B23R)	6.2e-159	110/110	100	(Goebel et al., 1990)
DIL	1 .	91		VAR-BSH	9.1e-31	46/49	93	(Shchelkunov et al., 1995)
	1	669 452		CPX host range VAR-I D6L (BSH:D8L)	1.1e-13	22/50	44	(Spehner er al., 1988)
	1	574		VAR-I BISR (BSH: BISR)	1.7e-11 1.2e-05	21/50	42	(Shchelkunov et al., 1995)
		574		VAC BISR (WR: BITR)	8.6c-05	22/73 22/73	30 30	(Shchelkunov et al., 1995)
	1	634		VAC C9L	0.00011	11/24	45	(Goebel et al., 1990) (Kotwal and Moss, 1988a)
		585		VAR-I GIR	0.00013	22/74	29	(Shcheikunov et ul., 1995)
	ļ	516		orf virus	0.0088	15/49	30	(Sullivan et al., 1995b)
	ı	153		VAR-I D7L (BSH:DIOL)	0.014	12/28	42	(Shchelkunov et al., 1995)
005R	10203	140	15.5	Growth factor (EGF				
	10625			receptor binding)				(Twardzik et al., 1985)
CIIR		142		VAC	2.9e-82	99/104	95	(Stroobant et al., 1985)
D2R		140		VAR-I (BSH:D4R)	3.6c-74	106/140		(Goebel et al., 1990) (Shehelkunov et al., 1995)
		138		CPX D5R	3.4e-95	101/114		(Safronov et al., 1995)
		169		human epiregulin	2.2c-14	29/78	37	D30783
				100 matches to growth factor	<0.10			
•				like sequences				
006L	11758	326	37.9	37.9k protein				(Variations to 1999)
CIOL	10778	331		VAC	1.7e-235	264/268	98	(Venkatesan et al., 1982) (Goebel et al., 1990)
201		331		CPX D6L	7.7e-235			(Safronov et al., 1996)
D5L		330 316		VAR-BSH (I: D3L)	3.6e-233	169/171		(Shchelkunov et al., 1995)
		316		VAR-J DIIL (BSH:DI4L) VAC C4L	1.7e-94	34/68	44	(Shchelkunov et al., 1995)
		315		CPX DI6L	1.8e-92 2.3e-92	30/68 31/68	54 45	(Goebel et al., 1990)
		82	•	Ectrometia 42K protein	1.2e-50	78/82	95	(Safronov et al., 1996)
		418		FPV BamHI ORFI	3.0e-11	13/41	31	(Senkevich et al., 1993a) (Tomley et al., 1988)
07R	12263		10.6	***				(10) 27 41., 1568)
U / IC	12538	91 242	10.6	28k virulence factor (f)				(Senkevich et al., 1993a)
	030	184		CPX D7R VAC-WR 21.7k protein	1.5c-51	42/47	89	(Safronov et al., 1996)
4R		242		VAR-I (BSH:D6R)	5.3c-51 3.7e-50	41/47 41/47	87	(Kotwal and Moss, 1988a)
	•	241		Ectromelia 28k secreted	3.7e-50	41/47	87 87	(Shchelkunov et al., 1995)
				virulence factor			٠.	(Senkevich et al., 1993a)
08L	13414	120	12.7	72 91				
708L	13414	120 126	13.7	13.7k protein				
	. 2034	138		VAR-BSH (1:DSL) Ectromelia 16k protein	1.9c-83	57/64	89	(Shchelkunov et al., 1995)
		124		CPX D8L	7.8e-81 3.2e-67	58/60 49/60	96	(Senkevich et al., 1993a)
		68		7.8k protein (VAC-WR)	1.3e-34	49/60 53/64	8 I 8 2	(Safronov et al., 1996) (Kotwal and Moss, 1988a)
205				•		30,07	J.	(2000 Bild WOS, 19882)
09L	13745	90	10.7	77k CPX hr protein (f1)				(Spehner et al., 1988)
1	13473	669		CPX host range gene	2.7e-46	43/52	82	(Safronov et al., 1996)
10L	14186	634 142	16.1	VAC C9L	1.7e-05	9/33	27	(Goebel et al., 1990)
	13758	669	10.1	77k CPX hr protein (f2) CPX host range gene	2 24 01	122/11-		(Spehner et al., 1988)
. 1		634		VAC C9L	2.2e-91 9.2e-21	133/142 26/63		(Safronov et al., 1996)
6L		452		VAR-I (BSH: D8L)	4.5c-13	27/29		(Goebel et al., 1990) (Shchelkunov et al., 1995)
ļ		150		VAC CI8L/B24R	1.3e-11	19/52		(Goebei et al., 1995)
į		439		AT ankyrin repeat protein	9.5e-07	23/59	38	(Zhang et al., 1992)
1		558		VAR-I BGR (BSH:B5R)	4.0e-05			(Shchelkunov et al., 1995)
ł				30 matches with ankyrin repeat containing proteins	2.7e-05 to			· · · - •
	14682	135	15.8	77k CPX hr protein (f3)	0.016			(B. ) (
IIL								(Spehner et al., 1988) .
1	14275	669		CPX host range gene	7.6e-80	54/64		
06L		669 452 90	10.3	CPX host range gene VAR-I (BSH: D8L) 77k CPX hr protein (f4)	7.6e-80 9.2c-78		84	(Safronov et al., 1996) (Shcheikunov et al., 1995)

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-30- GENOMIC SEQUENCE OF THE MVA STRAIN

STOP	ORF'	STAR	T AA	k Dar	name / (putative)				
Dell		STOP			(paramet)				
1					VAR-L(BSH: DRL)	2 2- 62	00105		
1.00	<b>'</b>	1	669	)					(Shchelkunov et al., 1995)
	ı	- 1			VAR-I D7L (BSH: D10L)				(Shehelkunov et al. 1988)
131L   15420   71   71   72   72   73   73   73   74   74   74   74   74		ŀ			C hotelinum NENII				(Goebel et al., 1990)
1305   1500		ļ			Caprings NINH protein				(Hutson et al., 1996)
013L   15420   15   15   15   15   15   15   15   1		i	895						
131L   15420   71   8.5   77k   CPX hp protein   15205   669   15205   669   15205   669   15205   669   15205   669   15205   669   15205   669   15205   669   15205   669   15205   669   15205   669   15205   669   15205   669   15205   669   15205		1			orf virus ank-like	0.0064			(Bull et al., 1996)
Del.   15205   669     CPX host range seas   15205   669     CPX host range seas   7.94-12   64/67   98   CS (Seftenor et al., 1988)   15205   669     CPX DIL.   147/31   42   CGPL DIL.   147/31   43   CGPL DIL.			6/3		rabbit fibroma 77.2k protei	0.0072			(Massung et al., 1992)
Doc	013L				77k CPX hr protein (f5)				(Spehner et al., 1988)
	D6L	1.5205			VAR (BSH: DRI)				(Safronov et al., 1996)
1.00					rabbit fibroma 77.2k proteir	0.0052			(Shchelkunov et al., 1995)
1-14		1			VAC CI7L/B23R				(Goebel et al., 1992)
									(Sullivan et al., 1995h)
					VAR BISK (WK: BITK)				(Goebel et al., 1990)
Column   C	014L	16205	109	13.1	•		7-1-0	**	
D9L	C9L	15876			VAC	3 973	100/100	3 100	(Kotwal and Moss, 1988a)
10	D01								(Goebel et al., 1990)
1515	<i>0</i> 9L .	1 .							(Shchelkunov et al. 1995)
1615   16496   64   11.2   75k   ank-like gene (T2)   75k   ank-like gene		Ì				3.7c-19			(Safronov et al., 1996)
CSPL			96	11.2	75k ank-like gene (f7)	0.021	5/16	31	(Massung et al., 1992)
1914	C9L	16496			VAC	4.0e-53	80/80	100	(Goebel et al. 1998a)
172		}				3.9e-25	48/80		(Safronov et al., 1990)
141		1							(Safronov et al., 1996)
O.65L   177   50   50   177   50   50   178   50   50   178   50   50   178   50   50   178   50   50   178   50   50   50   50   50   50   50   5		1							(Massung et al., 1996)
1759   1779   297   35.0   75k ank-like gene (13)   3.4e-208   291/294   98   (Kowal and Moss, 1988a) (Kowal and Moss, 1988a		1			CPX host range gene				(Schweizer and Neumann, 1995)
D7L				35.0	75k ank-like gene (f3)		• • • • •	32	(Kotwal and Moss 1988)
D7L	CYL	10800						98	(Goebel et al., 1990)
DRL	D7L								(Safronov et al., 1996)
DBL   452   CPX D9L   2,2e-16   23/61   37   (Safronov et al., 1986)   383   386   VAC C17L/B23R   2,9e-08   11/24   45   (Goebel et al., 1996)   383   CPX D9L   2022   Capripox virus   0.0085   13/58   22   (Goebel et al., 1995)   (Goebel et al., 1996)   (Goebel et al.		ľ							(Shchelkunov et al., 1995)
336	001				CPX D9L				(Safronov et al., 1988)
Sample   S	Dal	1						34	(Shchelkunov et al., 1995)
174									(Goebel et al., 1990)
202   Capripox virus   0.084   11/29   37   (Sullivan et al., 1995)		1	574						
171		1			Capripox virus				
CSL   17802   184		_j	374		VAR-1 B19R (BSH:B16R)	0.090	13/40	32	(Shchelkunov et al., 1995)
182				20.8	20.8k protein				(Kotwal and Moss, 1988a)
182		.,,,,,							(Goebel et al., 1990)
18859   150   18.0   host range protein									(Safronov et al., 1996)
C7L   18407   150			795		VAC H4L (RAP94)				
DILL   150				18.0					(Perkus et al., 1991)
185		10407							(Kotwal and Moss, 1988a)
197									(Shchelkunov et al., 1995)
170					Capripox virus ORF CF8a				(Schnitzlein and Tripathy, [991)
128   Myxoma virus ORF MF8   5.6e-13   16/43   37   (Jackson and Bults, 1992)		•			CPX D4L				(Safronov et al., 1989a)
1954							16/43	37	(Jackson and Bults, 1992)
Cold   19068   151	101	10541				J.4e-06	18/60	30	(Shchelkunov et al., 1995)
156	6L			18.2		76-101	1511151	100	(Kotwal and Moss, 1988a)
156	9L		156		VAR (BSH: DI2L)				(Goebel et al., 1990)
151						1.3c-96			(Safronov et al., 1996)
181									(Gershon and Black, 1989a)
149									(Upton et al., 1987)
149   VAC-WR K7R   0.40   8/13   61   (Kotwal and Moss, 1988a)   1988a)   1988a   19			149						
113   13.2   14k virulence factor, secreted protein (f)   117   VAC   2.6c-60   92/102   90   (Kotwal and Moss, 1988a)   117   VAR-BSH, virokine   6.6c-56   88/102   83   (Shchelkunov et al., 1995)   107   Rabbit fibroma virus   0.015   10/17   58   (Safronov et al., 1996)   (Massung et al., 1992)   107			149		VAC-WR K7R				(Kotwal and Moss, 1988a)
VIL	20L		113	13.2					
PIL 117 CPX PIL 7.3e-58 85/102 83 (Schecklunov et al., 1990) PIL 117 VAR-BSH, virokine 6.6e-56 88/102 86 (Safronov et al., 1995) PIL 20656 170 20.3 alpha-amanitin sensitive protein (Cotward and Moss, 1988a) PIL 20144 175 CPX PIL 3.0e-118 138/142 97 (Safronov et al., 1996) PIL 20144 175 VAC 6.1e-118 137/142 96 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1990) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1996) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1996) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1996) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1996) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1996) PIL 20144 175 VAR 9.7e-115 135/142 95 (Goebel et al., 1996)	11.	19084	117						(Kotwal and Moss. 1988h)
117   VAR-BSH, virokine   7.3e-38   85/102   83   (Shchelkunov et al., 1995)	- <b>-</b>								(Goebel et al., 1990)
107   Rabbit fibroma virus   0.015   10/17   58   (Massung et al., 1992)   121L   20656   170   20.3   alpha-amanitin sensitive   (Tamin et al., 1991)   (Kotwal and Moss, 1988a)   175   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1996)   (Yar and Moss, 1988a)   175   VAC   6.1e-118   137/142   96   (Goebel et al., 1996)   (Goebel et al., 1996)   (Safronov et al., 1996)   177   VAR   9.7e-115   135/142   95   (Shchelkunov et al., 1995)   180	IL		117						(Shchelkunov et al., 1995)
221L   20656   170   20.3   alpha-amanitia   sensitive			107		Rabbit fibroma virus				
175   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1996)     2L   175   VAC   G.1e-118   137/142   96   (Goebel et al., 1996)     2L   20981   98   11.0   33k host range gene (f)     22L   20985   284   VAC   1.8e-56   86/88   97   (Altenburger et al., 1989)     31L   66   VAC   2.3e-56   86/88   97   (Safronov et al., 1989)     31L   66   VAC   (Safronov et al., 1986)     32R   CPX MIL   2.3e-56   86/88   97   (Safronov et al., 1989)     33R   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1989)     34   CPX MIL   2.3e-56   86/88   97   (Safronov et al., 1989)     35   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     36   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1996)     37   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1996)     38   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     38   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1996)     38   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     38   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1995)     38   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     38   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     38   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     39   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     30   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     30   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     30   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     30   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     31   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     31   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     32   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     32   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     32   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     32   CPX P2L   3.0e-118   138/142   97   (Safronov et al., 1980)     33   CP			170	20.3					•
175		20144	175		protein				(Kotwal and Moss, 1988a)
2L 177 VAR 9.1c-118 137/142 96 (Goebel et al., 1990)  22L 20981 98 11.0 33k host range gene (f)  7L 20685 284 VAC 1.8c-56 86/88 97 (Altenburger et al., 1989)  32L 2088 98 11.0 33k host range gene (f)  4.8c-56 86/88 97 (Altenburger et al., 1989)  32L 20885 284 CPX MIL 2.3c-56 86/88 97 (Safronov et al., 1989)	2L								(Safronov et al., 1996)
22L 20981 98 11.0 33k host range gene (f)  (Gillard et al., 1986)  VAC 1.8c-56 86/88 97 (Altenburger et al., 1989)  1L 66 VAS 2.3c-56 86/88 97 (Safronov et al., 1996)									(Goebel et al., 1990)
7L     20685     284     VAC     1.8c-56     86/88     97     (Gillard et al., 1986)       284     CPX MIL     2.3c-56     86/88     97     (Safronov et al., 1996)       3L     66     VAC     2.3c-56     86/88     97     (Safronov et al., 1996)	22L	20981	98	110		,.,u-(1)	153/144	,,	(Girchetkunov et al., 1995)
284 CPX MIL 1.8c-36 86/88 97 (Altenburger et al., 1989) 1L 66 VAR 2.3c-56 86/88 97 (Safronov et al., 1996)				11.0		10000	06100	0.7	
			284		CPX MIL				(Altenburger et al., 1989)
2.0e-39 63/66 95 (Shchelkunov et al. 1995)	_		66 65		VAR	2.0e-39	63/66		(Shchelkunov et al., 1990)
65 human NOTCH 2 0.00036 17/41 41 (Kutsanis et al., 1995)					numan NOTCH 2				(Kutsanis et al., 1996)

ORF*	START	AAb	kDa¢	name /	(putative) / homologies*	BLAST <sup>d</sup> expect	BLAST'	HSS1 (%)	references
	erminal	regio						0 /	
023L	22296 21187	369	42.3		SPI-3, cell-cell mutation				(Boursnell et al., 1988)
K2L	-	369		VAC		1.2c-258	365/369	98	(Altenburger et al., 1989) (Goebel et al., 1990)]
C2L		373 373		CPX M2L VAR-BSH		1.2c-256 9.9e-249	331/337	0.5	(Safronov et al., 1996)
		373			virus H14-B	6.5e-244	321/337 312/337	73	(Shchelkunov et al., 1995) U67964
		386		HS plasm	iinogen activator	1.1e-35	30/68	44	(Loskutoff 'et al., 1987)
		58		inhibitor CPX SPI		8.2e-33	57/58	98	•
		369		Myxoma 1	virus MAPI gene	7.3e-32	33/131	25	gi:1168082 (Upton et al., 1990a)
		397 397			otease nexin	1.5e-29	31/67	46	(Vassalli et al., 1993)
				promoting		8.7e-27	30/65	46	A03911
		320 417		Swinepox	SPI like protein	3.6e-21	20/70	28	(Massung et al., 1993)
		383			ypsin, human croid-binding	2,2c-20 9.0c-20	26/66	39	(Ciliberto et al., 1985) (Seralini et al., 1989)
		200		protein (	rabbit)				
		390		squamous antigen	cell carcinoma	1.9c-17			(Schneider et al., 1995)
024L	22612	88	10.5		stance, eIF-2a				(Beattie et al., 1991)
	22346	88		homolog		26- 6-	00/00	100	(Davies et al., 1992)
K3L		88		CPX M3L VAC		2.6e-61 1.4e-60	88/88 87/88	100 98	(Safronov et al., 1996) (Goebel et al., 1990)
C3L	<i>,</i> .	88 .		VAR-I		1.0e-52	73/88	82	(Shchelkunov et al., 1995)
		86		SPV C8 p	rotein initiation factor 2	4.1e-22 1.2e-08/	20/44	45 -	(Massung et al., 1993)
				family		0.45			•
025L	23938 22664	424	48.9	phosphol protein	ipase D-like				(Cao et al., 1997)
K4L		424		VAC		1.5e-306	423/424		(Goebel et al., 1990)
		424 437		CPX M4L human HI	I-K4	2.1e-303 2.8e-135	416/424 53/95	98 55	(Safronov et al., 1996)
		372		D. discoid	cum	2.5e-91	28/47	59	U60644 (Giorda et al., 1989)
		516		C. elegans	•	6.6c-89	31/61	50	gi: 2435624
		2327 635		C. elegans C. elegans		2.8c-52 1.1e-24	36/60 19/53	60 35	gi: 2291241 (Wilson <i>et al.</i> , 1994)
		377		FPV majo	r envelope protein	2.9e-23	19/61	31	(Calvert et al., 1992)
		371 378		Orf virus	rirus env protein env protein B2L	3.6e-22 1.2e-21	18/51	35 29	U43549
MC021L		388		MCV subt	ype I env protein -	3.2e-21	20/63	3 l	(Sullivan <i>et al.</i> , 1994) (Senkevich <i>et al.</i> , 1997)
CI7L		372 372		VAR-BSH VAC FI3L	•	4.6e-19 4.9e-17	15/52 15/52	28 28	(Shchelkunov et al., 1995) (Goebel et al., 1990)
026L	124478	170	19.1	_	pholipase-like	- •			•
	23966			protein	(f1)				(Upton & Buller, unpub.)
1	1	276 277		CPX M5L Ectromelia	virus HI4-E	2.6c-110 2.7c-109	161/170		(Safronov et al., 1996)
KSL	1	136		VAC	us 1117°E	5.5c-69	160/170		X94355 U67964 (Goebel et al., 1990)
1		134 313		VAC-WR	hornholiness	8.3e-63	98/101	97	(Boursnell et al., 1988)
1	[	323		homolog	hospholipase	3.3e-35 1.2e-13	35/105 30/94	33 31	U67963 297050
1	1		•	poss. oxide	oreductase M.				
1 .	1	324		Lysophosp	sum holipase isolog	3.1c-5	13/58	22	U95973
1		313		A. thalian		0.047	13/30	43	U32747
0277	24604		7.0	lysophosph	olipase L2				
027L	24694 24500	64	7.0	lysophos protein	pholipase-like (f2)	-			(Upton & Buller, unpub.)
K6L	1	81		VAC	•	5.3e-42	63/63	100	(Boursnell et al., 1988)
1	l	276 277		CPX M5L Ectromelia	virus H14-E	2.4c-36 2.4c-36	57/58 57/58	98 98	(Safronov et al., 1996) U67964
	1	313		HS lyopho	ospholipase homolog	9.1c-23	34/53	64	U67963
]	]	323		hyp. oxido	reductase M.	9.9e-14	22/54	40	Z97050
	1	530		tuberculo: dihydrote:	iis stosterone/androsta	7.0e-05	6/17	35	A48633
					DP-glucuronosyl-		•		•
central	_conser	ved	region:						1
028R	24864	149	17.5		otein				(Goebel et al., 1990)
K7R	25313	149 161		VAC CPX M6R		6.1e-105	149/149		(Goebel et al., 1990)
C4R		149		VAR		1.6e-101 4.9e-101	144/149 143/149		(Safronov et al., 1996) (Shchelkunov et al., 1995)
		236		Swinepox	(sc76)	0.00017	19/49	95	(Massung et al., 1993)
029L	26046	222	25.9	25.9k pr	otein				(Roseman and Slabaugh, 1990)
FIL	25378	226		VAC		2.7e-158	208/211		(Goebel et al., 1990)
C5L		238 251		CPX GIL VAR-I		7.0e-148 6.6e-147	166/189 184/200		(Safronov et al., 1996) (Shchelkunov et al., 1995)
030L	26501		16.2						•
	26501 26058	147	16.2	dUTPase					(Roseman and Slabaugh, 1990) (Roseman et al., 1996)
F2L		147		VAC		2.9e-102	147/147		(Goebel et al., 1990)
C6L		147 147		CPX G2L V A R		8.2c-100 1.1c-97	144/147 142/147		(Safronov <i>et al.</i> , 1996) (Shchelkunov <i>et al.</i> , 1995)
		164		human dU	TPase	4.1c-61	49/69	71	(Ladner et al., 1996)

-32- GENOMIC SEQUENCE OF THE MVA STRAIN

ORF'	STAD	T 4.4	b 1:12s					•
	STOP	TAA		name / (putative) function / homologies*	BLAST expect	BLAST <sup>c</sup>	HSS	references
1611	terminal	regi		Swinepox virus	8.0e-56			
		15		orf virus avian adenovirus	1.5c-49	45/69	61 65	(Massung et al., -1993) (Mercer et al., 1989)
		11	-	FIV pol polyprotein	6.6e-49 1.5e-26		57 41	(Akopian et al., 1992) (Talbott et al., 1989)
				dUTPase pyrophosphatase family	>4.2e-06		••	(1210011 21 21., 1989)
031L	27955 26525		5 55.3	kelch-like protein				(Senkevich et al., 1993b)
F3L		480 483		VAC	0.0	292/29	4 99	(Roseman and Slabaugh, 1990) (Goebel et al., 1990)
C7L		179	)	CPX G3L V A R-1	0.0° 1.9e-124	287/29: 166/17		(Safronov et al., 1996)
		500 564		Swinepox virus protein C13 VAC A55R	4.4c-46	39/133	29	(Shchelkunov et al., 1995) (Massung et al., 1993)
		689		kelch protein D.melanogaste	2.8e-21 r 5.3e-18	17/51 21/65	33 32	(Goebel et al., 1990) (Xue and Cooley, 1993)
		512 512	:	CPX D18L VAC C2L	1.4e-16 1.6e-16	15/33 15/33	45	(Safronov et al., 1996)
		625 624		T27E9.4 C. elegans	3.7c-14	15/59	45 25	(Goebel et al., 1990) Z82059
		817		human KIAA0132 protein R09A8.3 (C. elegans)	1.9c-13 1.1c-12	13/60 17/45	21 37	D50922 o.k (Wilson <i>et al.</i> , 1994)
		611 530		C47D12.7 (C. elegans) Swinepox virus	2.4c-12 3.0e-09	22/91 14/58	24	(Wilson et al., 1994)
		589 <b>5</b> 21		M M actin binding protein CPX C3L	1.9e-09	. 18/88	24 20	(Massung <i>et al</i> ., 1993) U65079
		509 202		Myxoma virus MT-9	1.2e-08 2.5e-08	15/37 17/58	40 29	(Safronov <i>et al.</i> , 1996) (Upton <i>et al.</i> , 1990a)
				Murine IAP-promoted placenta (MIPP) expressed	4.3e-08	17/56	30	(Chang-Yeh et al., 1991)
		326 559		protein A. thaliana hyp. protein	3.9e-06 9.0e-6	22/80	27	299708
		916 172		Ectromelia virus p65	0.00016	12/31 13/42	38 30	(Senkevich et al., 1993b) (Way et al., 1995)
		1,72		B-scruin (L. polyphemus) VAR-I J8R (BSH: J6R)	810.0	15/36	41	(Shehelkunov et al., 1995)
032L	28925	319	37.0	ribonucleotide reductase				
	27966	319		(small subunit) CPX G4L				(Slabaugh et al., 1988) (Roseman and Slabaugh, 1990)
F4L		319		VAC	2.3e-231 3.5e-231	317/319 317/319		(Safronov et al., 1996) (Goebel et al., 1990)
C8L		333		VAR-BSH ribonucleotide reductase	4.1e-228	313/319		(Shchelkunov et al., 1995)
				family	>2.2e-10			
033L	29250 28957	97	11.1	36.5k major membrane				(Roseman and Slabaugh, 1990)
C9L	20937	348		protein precursor (f1) VAR-BSH	1.9e-36	51/53	96	
FŚL		323 321		CPX G5L VAC	2.4c-19	47/77	61	(Shchelkunov et al., 1995) (Safronov et al., 1996)
1		1584		non-receptor tyrosin kinase	3.3e-19 0.00038	42/70 15/35	60 42	(Goebel et al., 1990) (Tan and Spudich, 1990)
034L	29875	218	24.8	(Dictyostelium discoideum) 36.5k major membrane				
1	29219	323		protein precursor (f2) CPX GSL	8.2e-155	215/217		(Roseman and Slabaugh, 1990)
F5L C9L		321 348		VAC VAR-BSH	6.4e-155		99	(Safronov et al., 1996) (Goebel et al., 1990)
035L	 			•	6.8c-141	186/210	88	(Shchelkunov et al., 1995)
F6L	30129 29905	74 74	8.6	8.6k protein VAC	5.5e-47	74/74	100	(Roseman and Slabaugh, 1990)
CIOL		72		VAR	2.3c-38	62/70	88	(Goebel et al., 1990) (Shehelkunov et al., 1995)
036L CIIL	30387 30145	80 79	9.4	9.4k protein				(Roseman and Slabaugh, 1990)
F7L	50.15	92		VAR VAC	2.9e-44 1.9e-43	34/43 65/65	79 100	(Shchelkunov et al., 1995)
037L	30731	65	7.9	7.9k protein		•	-	(Goebel et al., 1990)
F8L Cl2L	30534	65 65		VAC VAR-I	5.1c-43		96	(Roseman and Slabaugh, 1990) (Goebel et al., 1990),
038L	31429 /	212	23.8		3.1e-41	61/65	93	(Shehelkunov et al., 1995)
F9L	30791	212		VAC	7.1e-148	212/212	001	(Roseman and Slabaugh, 1990) (Goebel et al., 1990),
CI3L		212 215		VAR Swinepox virus	1.2e-144 8.1e-72	207/212	97	(Shehelkunov et al., 1995)
MC016L		213 225		MCV subtype I Orf virus	2.8e-62	71/152	4 l 46	(Massung et al., 1993) (Senkevich et al., 1996)
		243		FPV protein FP2	5.1e-39 2.8e-17		32 44	(Mercer et al., 1995) (Binns et al., 1988)
		243 250		MCV subtype 1 MC069R VAC LIR	7.7e-12 1.1e-07	23/58	39	(Senkevich et al., 1996)
		250		VAR MIR	1.1e-07		34 34	(Goebel et al., 1990). (Shchelkunov et al., 1995)
039L	32735 31416	439	52.1	scrine/threonine protein				(Lin and Broyles, 1994)
FIOL	2.710	439		kinase 2 VAC	0.0	429/439	<del>9</del> 7	(Wang and Shuman, 1995)
C14L		439 440		VAR-BSH Swinepox virus	0.0 2.2e-233	424/439	96	(Goebel et al., 1990), (Shchelkunov et al., 1995)
MC017L		443 498		MCV subtype 1 orf virus	2.3e-198	151/214 178/282	53	(Massung et al., 1993) (Senkevich et al., 1996)
040L	33012	84	0.6		2.2e-162	198/366	14	(Mercer et al., 1995)
CISL	32758	354	9.6	39.7k protein (ft) VAR	6.6e-27	50/64 7	8	(Shehalkuman
FIIL 041L	33771	354 100	11.4		9.1c-27			(Shchelkunov et al., 1995) (Goebel et al., 1990)
				p-5tern (12)				

				·				
ORF'	START	' AA	kDat	name / (putative)	BLAST	BLAST		references
left to	erminal	regio	n:	function / homologies	expect	AA Id	(%)	
FIIL	33469	354		VAC	3.8e-62	95/95	100	(0-1)
CISL	_	354		VAR	8.8e-58	90/95	94	(Goebel et al., 1990), (Shehelkunov et al., 1995)
042L	35721	635	73.1	73.1k protein				(======================================
FIZL	33814	635	73.1	73.1k protein VAC	0.0	629/635	99	(Carba)
C16L		635		VAR-I	0.0	607/635		(Goebel et al., 1990), (Shchelkunov et al., 1995)
MC019L		352 663		Myxoma virus MCV subtype I	3.6c-84	28/66	42	U43549
MCOISE	•	640		orf virus	4.0c-60 4.8c-39	29/82 19/61	35 31	(Senkevich et al., 1996)
		630		FPV F12 homolog	2.3e-15	19/67	28	U34774 (Ogawa et al., 1993)
043L	36866	372	41.8	37k malan EEN				·
*****	35748	3,2	41.0	37k major EEV antigen IMCBH sensitive protein				(Hirt et al., 1986)
E121		272		palmitylprotein				(Schmutz et al., 1991) (Grosenbach et al., 1997)
<i>F13L</i> C17L		372 372		VAC VAR-BSH	2.1e-268 8.9e-265			(Gocbel et al., 1990)
		371		Myxoma virus	2.5e-115	110/200		(Shchelkunov <i>er al.</i> , 1995) U43549
MC021L		378 388		orf virus	7.6e-108	83/194	42	(Sullivan et al., 1994)
MCOZIC	1	377		MCV subtype 1 FPV major env protein	6.1e-98 2.8e-88	44/113 47/112	38 41	(Senkevich et al., 1996)
		251		pigeonpox virus	1.8e-62	47/112	41	(Calvert et al., 1992) S27933
		424		CPX M4L VAC K4L	2.1e-18	16/52	30	(Safronov et al., 1996)
		372		D. discoideum	1.7c-17 1.4c-16	14/35 28/84	40 33	(Goebel et al., 1990)
		437		HU-K4 (homo sapiens)	1.5e-11	25/94	26	(Giorda <i>et al.</i> , 1989) U60644
044L	37105	73	0 2	9 7te				
FI4L	36884	73	8.3	8.3k protein VAC	2.3c-44	72/73	98	(Goshel et al. 1000)
C18L		73		VAR	2.1c-35	57/73	78	(Goebei et al., 1990) (Shchelkunov et al., 1995)
045L	378533	158	18.6	18.6k protein		-		
FISL	37377	158	. a. 0	18.6k protein VAC	2.3c-112	157/158	99	(Goebel et al., 1990),
CI9L		161		VAR	1.46-107	150/153		(Shchelkunov et al., 1995)
MC025L		148 148		MCV subtype I Myxoma virus	3.5c-54	52/113	46	(Senkevich et al., 1996)
		174		Myxonia Viius	5.4e-50	48/112	42	U43549
046L	38555	23!	26.5	26.5k protein		•		
<i>F16L</i> C20L	37860	231 231		VAC VAR	3.3e-159 5.6e-157	227/231		(Goebel et al., 1990),
		209		Myxoma virus	8.3c-48	222/231 26/58	96 44	(Slichelkunov et al., 1995) U43549
MC029L		230		MCV subtype 1	6.9c-45	16/61	26	(Senkevich et al., 1996)
047R	38619	101	11.3	11k DNA binding				(Parchalas as at 1996)
	38924			phosphoprotein				(Bertholet et al., 1985) (Kao and Bauer, 1987)
F17R C21R		101 101		VAC	3.0€-69	100/101	99	(Goebel et al., 1990)
Carr		102		VAR MYX	9.7e-67 6.6e-26	99/101 45/92	98 98	(Shchelkunov et al., 1995) U43549
MC030R		92 .		MCV subtype 1	1.5e-20	33/53	48	(Senkevich et al., 1997)
		46		orf virus	1.3e-06	16/29	62	(Mercer et al., 1995)
048L	40360	479	55.6	poly(A) polymerase				(Gershon et al., 1991)
EIL	38921	479		catalytic subunit				
EIL		479		VAC VAR-I	0.0 0.0	478/479 472/479		(Goebel et al., 1990),
MC031L		470		MCV subtype 1	1.5c-177	114/173		(Shchelkunov et al., 1995) (Senkevich et al., 1997)
049L	42570	737	85.9	85.9k protein				
E2L	40357	737	05.5	VAC.	0.0	735/737	99	(Ahn et al., 1990a) (Goebel et al., 1990),
E2L MC032L		737		VAR-I	0.0	731/737	99	(Shchelkunov et al., 1995)
MCUJZL		748		MCV subtype I	8.3c-127	59/198	29	(Senkevich et al., 1997)
050L	43269	190	21.5	dsRNA dependent PK				(Chang et al., 1992)
E3L	42697	190		Inhibitor, host range VAC	1 4- 100	10011		(Chang et al., 1995b)
E3L		192		VAR-BSH	1.4e-129 8.6c-126	188/190		(Gocbel et al., 1990), (Shchelkunov et al., 1995)
		1175		dsRNA specific ADA (rat)	7.2c-12			(O'Conneil et al., 1995)
		1226 551		dsRNA specific ADA (human) human protein kinase p68	2.8c-09		44	(Kim er al., 1994)
		•		INF inducible kinase family	3.8c-05 >0.00099	22/42	52	(Meurs et al., 1990)
051L	44 1 0 3	259	20.0	·				
VJ11	43324	439	29.8	RNA polymerase subunit rpo30, VITF-1				(Ahn et al., 1990a)
E4L		259		VAC	1.6c-182	258/259	99	(Broyles and Pennington, 1990) (Goebel et al., 1990)
E4L MC034L		259 444		VAR-BSH MCV subtype I	3.2e-180	255/259	98	(Sheheikunov et al., 1995)
		39		orf virus	1.2c-84 6.7c-10	107/171 21/39		(Senkevich et al., 1996) (Mercer et al., 1995)
		243		African swine fever virus	0.00034			(Vydelingum et al., 1993)
				TFIIS family	<0.0096			-,
	44180	331	39.1	39.1k protein				(Goebel et al., 1990)
ESR ESR	45175	331 341		VAC	1.2e-235	329/331	99	(Goebel et al., 1990)
<b>₩</b>		332		VAR Talerapox	3.1c-223 7.1c-225	312/331 300/314	94	(Shchelkunov et al., 1995)
		329		Camelpox	1.46-221	206/220		(Douglas and Dumbell, 1996) (Douglas and Dumbell, 1996)
		319 256		Cowpox	1.5e-202	271/303	89	(Douglas and Dumbell, 1996)
MC038R		276		Ectromelia MCV subtype 1	3.8c-153 8.3c-109	218/245 94/152	88	(Douglas and Dumbell, 1996) (Senkevich et al., 1997)
053R	45312	567	66.7	**				
				66.7k pratein				(Goebel et al., 1990)

-34-GENOMIC SEQUENCE OF THE MVA STRAIN

ORF.	STAF	T AA'	kDa	(paratio)	BLAST		* HSS	references
left	termina		on:	function / homologies	expect	AA id	(%)	
E6R MC037	R	567 565		VAR MCV subtype 1	0.0 7.2e-247	555/56 258/45		(Shchelkunov et al., 1995) (Senkevich et al., 1997)
054R <i>E7R</i> E7R	47082 47582	166 166 60	19.5	17k myristylprotein VAC VAR-I (BSH: E6.5R)	9.7e-116 2.7e-36	166/166 53/60	5 100 88	(Martin et al., 1997) (Goebel et al., 1990) (Shehelkunov et al., 1995)
055R E8R E8R MC038I	47695 48516 R	273 273 273 276	31.9	31.9k protein VAC VAR MCV subtype !	4.5e-195 9.9e-192 8.3e-109	266/273		(Earl et al., 1986) (Goebel et al., 1990) (Shchelkunov et al., 1993a), (Senkevich et al., 1997)
056L E9L E9L MC039L	51543 48523	1006 1005 1005 1008 988 1004 964		DNA polymerase VAC VAR BSH Orf virus FPV MCV subtype ! C. biennis poxvirus DNA polymerase family	0.0 0.0 0.0 0.0 0.0 2.6e-77 >6.0e-06	1005/10 06 598/608 199/388 179/294 175/297 28/82	98 <sup>-</sup> 51 60 58	(Earl et al., 1986) (Goebel et al., 1990), (Shchelkunov et al., 1995) (Mercer et al., 1996) (Binns et al., 1987) (Senkevich et al., 1997) (Mustafa and Yuen, 1991)
057 R <i>E10R</i> E10R MC040R	51575 51862	95 95 95 101	10.9	10.9k protein VAC VAR MCV subtype	1.2e-65 3.1e-64 5.2e-44	93/95 90/95 58/95	97 100 94	(Goebel et al., 1990) (Goebel et al., 1990) (Shchelkunov et al., 1993a) (Senkevich et al., 1997)
058L E11L E11L MC041L	52246 51857	129 129 129 132	14.9	14.9k protein VAC VAR MCV subtype 1	3.3e-89 4.2e-87 1.8e-30	129/129 125/129 31/96		(Goebel et al., 1990) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1997)
059L OIL QIL MC042L		152 666 666 783	17.6	77.6k protein (f1) VAC VAR-BSH MCV subtype ! leu zipper, bipartite nuclear targeting sequence	6.9e-101 3.4e-92 1.5e-22	151/152 137/152 39/105		(Goebel et al., 1990) (Goebel et al., 1990), (Shchelkunov et al., 1995) (Senkevich et al., 1997) (Goebel et al., 1990)
060L O/L QIL MC042L	54189 52972	405 666 666 783	47.4	77.6k protein (f2) VAC VAR-I MCV subtype !	5.8e-277 1.7e-269 2.7e-51	399/400 383/400 38/104		(Goebel et al., 1990) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1997)
061L O2L Q2L	54555 54229	108 108 106	12.4	glutaredoxin 1  VAC  VAR human glutaredoxin glutaredoxin family	2.0e-74 4.9e-72 3.2e-31 >9.0e-05	108/108 104/108 49/106		(Ahn and Moss. 1992a) (Johnson et al., 1991) (Goebel et al., 1990) (Shcheikunov et al., 1995) (Fernando et al., 1994)
062L // L KIL MC044L	55639 54701	312 312 312 310 1451	35.9	35.9k protein VAC VAR-BSH MCV subtype   transcription initiation protein (S. cerevisiae)	4.7e-208 4.8e-205 3.8e-110 0.029	310/312 305/312 163/307 10/28	97	(Schmitt and Stunnenberg, 1988) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1996) (Hansen et al., 1996)
063 L 12 L K2 L MC045 L	55867 55646	73 73 73 72 887	8.5	8.5k protein VAC VAR MCV subtype 1 hypothetical yeast protein	5.5e-50 5.5e-50 3.5e-18 8.1e-05	73/73 73/73 20/33 9/24	100 100 60 37	(Schmitt and Stunnenberg, 1988) (Goebel et al., 1990) (Shehelkunov et al., 1995) (Senkevich et al., 1996) \$48422
064L 13L K3L MC046L	56677 55868	269 269 269 288 209	30.0	DNA binding phospho- protein (F4L interacting) VAC VAR MCV subtype 1 FPV 13 protein	2.1c-173 2.5c-172 9.6c-66 8.4c-35			(Schmitt and Stunnenberg, 1988) (Davis and Mathews, 1993) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1996) A48563
065L 141. K4L	59075 56760	77 l 77 l 77 l	87.8	ribonucleotide reductase (large subunit) VAC VAR ribonucleotide red, family	0.0 0.0 >1.8e-05	771/771 761/771	100	(Schmitt and Stunnenberg, 1988) (Tengelsen et al., 1988) (Goebel et al., 1990) (Shchelkunov et al., 1995)
066L /5L KSL MC047L	59342 59103	79 79 79 82 81 321	8.8	8.8k protein VAC VAR MCV subtype   FPV 9.1k protein formate dep. nitrit reductase protein (H. influenzae) permease (b. subtilis)	6.3c-49 1.2c-47 2.6c-17 1.4c-12 0.00022	76/79 27/73 13/38 7/18	96 36 34 38	(Schmitt and Stunnenberg, 1988) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1996) (Binns et al., 1988) (Fleischmann et al., 1995)
	60509 59361		43.5	43.5k protein VAC VAC VAR MCV subtype 1	8.6e-268 3.1e-267 2.1e-99	382/382 380/382	100 99	gi:2415386 (Schmitt and Stunnenberg, 1988) (Goebel et al., 1990) (Shchelkkunov et al., 1995) (Senkevich et al., 1996)

ORF:	STOP		kDa	name / (putative) function / homologies*	BLAST expect		" HSS	
_ieit	ret mina	1 reg 390		FPV 16 protein				
		3,0		mitochondrial energy transfer proteins signature	1.4e-86	50/136	36	E48563, P12925 (Goebel et al., 1990)
068L <i>17L</i>	61773 60502	423 423		core proteia, topoisomerase II VAC	2.2	:		(Schmitt and Stunnenberg, 198 (Kane and Shuman, 1993)
K7L .		423		VAR	0.0 1.5e-306	420/423 419/423		(Goebel et al., 1990)
MC049L		515		MCV subtype I	1.9e-199			(Shchelkunov et al., 1995) (Senkevich et al., 1996)
		421 464		FPV 17 protein Amsacta moorei poxvirus	8.1e-180 3.2e-14			F48563 (Hall and Moyer, 1991)
069R	61776 63809	676	77.6	NPH-II, NTPase, RNA helicase				(Shuman, 1992), (Koonin and Senkevich, 1992)
<i>isr</i> Kbr		676 676		VAC VAR	0.0 0.0	674/676		(Goedel & al., 1990)
MC050R	1	684		MCV subtype 1	7.6e-227	665/676 144/272		(Schelkunov et al., 1995) (Schkevich et al., 1997)
		682		FPV virus 18FPV 61 matches mainly to RNA helicase family	4.2e-206 <0.38	98/178	55	(Binns et al., 1988)
070L	65588	591	68.0	68k protein				(Schmitt and Stunnenberg, 1988
<i>GIL</i> HIL	63813	591 591		VAC VAR-1	0.0	590/591		(Goedel et at., 1990)
MC056L		593	-	MCV subtype 1	0.0 1.2e-217	582/591 183/361		(Shchelkunov et al., 1995) (Senkevich et al., 1997)
)71L	65020	341	12.0	FPV	9.4e-75	45/101		H48563
	65920 65585	111	12.8	12.8k protein				(Schmitt and Stunnenberg, 1988 (Meis and Condit, 1991)
<i>G3L</i> K3L		111		VAC VAR	7.6e-74	111/111		(Goebel et al., 1990)
MC057L		108		MCV subtype 1	2.4e-71 0.00012	108/111 15/45	97 33	(Shchelkunov et al., 1995) (Senkevich et al., 1997)
72R	65914	220	25.8	IBT-dependent protein				(Meis and Condit, 1991)
<i>32R</i> H2R	66576	220 220		VAC VAR	1.9e-155 1.1e-151	220/220	100	(Goebel et al., 1990)
1C058R		246		MCV subtype 1	2.7e-36	214/220 42/135	97 31	(Shchelkunov et al., 1995) (Senkevich et al., 1997)
73L	66920 66546	124	14.0	glutaredoxin 2 membrane protein				(Gvakharia et al., 1996)
14L 14L		124		VAR	4.0e-83	123/124	99	(Jensen et al., 1996) (Shchelkunov et al., 1995)
4C059L		124 126		VAC MCV subtype I	7.5e-83 1.1e-21	123/124 21/51	99 41	(Goebel et al., 1990) (Senkevich et al., 1997)
74R 55R	66923 68227	434 434	49.9	49.8k protein				(Goebel et al., 1990)
15R		434		VAR	1.6e-305 1.9e-299	432/434 423/434	99 97	(Goehel et al., 1990) (Shchelkunov et al., 1995)
AC60R		437 1300		MCV subtype 1 HS CG1 protein	1.0c-55 0.015	56/119 22/82	47 26	(Senkevich et al., 1997) (Print et al., 1994)
75R	68235 68426	63	7.3	RNA polymerase subunit				(Amegadzie et al., 1992),
75.5R 15.5R		63 63		VAC . VAR	1.1e-40	63/63	100	(Meis and Condit, 1991) (Goebel et al., 1990)
1C061R		63 ·		MCV subtype 1	1.1c-39 9.3e-27	61/63 41/63	96 65	(Shchelkunov et al., 1995)
				35 matches mainly to RNA polymerases	<0.54	11705	03	(Senkevich et al., 1997)
	68428 68925	165 165	19.0	18.9k protein VAC	3.9 117	1601155		(Goebel et al., 1990)
6R		165		VAR	3.8e-116 1.5e-116	162/165 164/165	98 99	(Goebel et al., 1990) (Shchelkunov et al., 1995)
C062R		195		MCV subtype 1	3.0e-32	27/57	47 -	(Senkevich et al., 1997)
	70005 68890	371 371	42.0	42.0k protein VAC	5 24 255	270/221	00	(Schmitt and Stunnenberg, 1988)
7L		37 I		VAR	5.2e-255 7.1e-255	370/371 S	99	(Goebel et al., 1990) (Shchelkunov et al., 1995)
C065L	<b>3005</b> -	402		MCV subtype I	2.0e-109			(Senkevich et al., 1997)
	70036 <sub>.</sub> 70818	260	29.9	VLTF-1, late transcription factor				(Keck et al., 1990) (Wright et al., 1991)
8 <i>R</i> Br		260 260		VAC VAR-I	8.6-184	259/260	99	(Goebel et al., 1990)
C067R		260		MCV subtype 1	3.1e-183 8.5e-136	258/260 185/260	99 71	(Shchelkunov et al., 1995) (Senkevich et al., 1997)
	•00	260		FPV virus FPO	3.3e-129	175/250	67	(Binns et al., 1988)
9R '	70838 71860	340 340	38.9	37k myristylprotein VAC	3.7e-237	317/319		(Martin et al., 1997)
9R C068R		340		VAR	9.1c-236	315/319		(Goebel et al., 1990) (Shchelkunov et al., 1995)
CUGOR		342 336		MCV subtype I FPV virus FPI		59/127 59/124	46	(Senkevich et al., 1997) (Binns et al., 1988)
	71861 72613	250	27.3	25k myristylprotein				(Franke es al., 1990)
l R		250		IMV virion protein VAC	1.8c-175	250/250		(Martin et al., 1997)
1 R C069R		250		VAR .	6.4c-170	249/250	99	(Goebel et al., 1990) (Shcheikunov et al., 1995)
COOTE		243 243		MCV subtype 1 FPV virus FP2	6.5e-103	145/243	59	(Senkevich et al., 1997)
		212		VAC F9L		128/243 20/58	52 34	(Binns et al., 1988) (Goebel et al., 1990)
		212		VAR CI3L		20/58	34	(Shchelkunov et al., 1995)

-36-GENOMIC SEQUENCE OF THE MVA STRAIN

ORF	STAI	RT AA	* kDa	s nume / (putative)				
left	STOR		ion:	function / homologies	BLAST expec		HSS(	references
		213	3	MCV subtype I MC016L	1.6e-0.	7 12/52		
		215	5	Swinepox	3.3c-0.		22 29	(Senkevich et al., 1997) (Massung et al., 1993)
081F L2R			10.3	10.3k protein VAC				(Plucienniczak et al., 1985)
M2R		87		VAR	3.9e-57 4.0c-56		100 97	(Goebel et al., 1990)
MC07	OK	93 504	ļ	MCV subtype I Na* dependent phosphate	0.064	18/80	22	(Shchelkunov et al., 1995) (Senkevich et al., 1997)
				transporter C. elegans	6.9e-05	10/39	25	(Wilson et al., 1994)
		233 233		ATPase subunit T. cruzi Ca <sup>2*</sup> channel rat	0.013	16/44	36	U38184
		223 155		Ca2+ channel mouse	5.2c+0.2 7.1c+0.2		24 24	(Dubel et al., 1992)
		123	y	ABC transporter yeast	0.40	12/40	30	(Coppola et al., 1994) X97560
082L <i>L3L</i>	73950 72898			40.6k protein VAC				(Plucienniczak et al., 1985)
M3L		349		VAR	2.2e-25 1.5e-24			(Goedel et al., 1990)
MC07:	2L	310 301		MCV subtype 1 FPV F4 protein	1.5c-88	64/136	47	(Shchelkunov et al., 1995) (Senkevich et al., 1997)
083R	72076		•••		1.1e-80	58/134	43	(Binns et al., 1988)
	73975 74730		28.5	core protein VP8 DNA/RNA binding prote				(Yang and Bauer, 1988)
<i>L4R</i> M4R		251 251		VAC	5.6e-170	251/251	100	(Baylis and Smith, 1997)
MC073	3R	254		VAR MCV subtype 1	3.7-169 1.7e-76	250/251	99	(Shcheikunov et al., 1995)
	•	253		FPV virus FP5	6.4e-55	36/59 29/57	6 ! 50	(Senkevich et al., 1997) (Binns et al., 1988)
084R	74740	128	15.1	15.1k protein				1700)
<i>L5 R</i> M5R	75126	128 128		VAC 14.0k protein VAR	2.9e-89	127/128	99	(Goebel et al., 1990)
		129		FPV FP6	2.0-87 8.1e-16	125/128 19/45	97 42	(Shchelkunov et al., 1995)
MC074	K	146 152		MCV subtype ! melatonin receptor D. rerio	0.073	10/18	55	(Drillien et al., 1987) (Senkevich et al., 1997)
085R	75083				0.44	15/66	222	(Reppert et al., 1995)
JIR	75544	153 153	17.9	dimeric virion protein VAC	6.0- 102	150		(Holzer & Falkner, unpubl.)
LIR		159 147		VAR-I	6.0e-103 1.4e-101	152/153 149/153	99 97	(Goebel et al., 1990) (Shchelkunov et al., 1995)
		148		capripox CF7 myxoma MF7	6.5e-54 4.8e-51	53/90 54/93	58	(Ocishon and Black, 1989b)
MC075	R	183 148		MCV subtype ( FPV FP7	1.9c-47	47/93	58 50	(Jackson and Bults, 1992) (Senkevich et al., 1997)
	•				1.3e-35	37/84	44	(Drillien et al., 1987)
086R	75560 76093	177	20.0	thymidine kinase				(Hruby and Ball, 1982)
J2R L2R		177		VAC	5.7e-125	175/177	98	(Weir and Moss, 1983)
LZK		177		VAR 38 matches mainly to	2.7c-122	170/177	96	(Goebel et al., 1990) (Shehelkunov et al., 1995)
				thymidine kinase family	<0.18			, ,,,,
087R	76159	333	38.9	poly(A) polymerase su.				
J3R	77160	333		2'methyl transferase VAC				(Gershon et al., 1991) (Gershon and Moss, 1993)
L3R		333		VAR-BSH	8.7e-136 9.8e-233	330/333 326/333	99 97	(Goebel <i>et al.</i> , 1990)
MC076R	t	338 343		myxoma MCV subtype t	5.7e-288	247/333	74	(Shchelkunov et al., 1995) (Jackson and Bults, 1990)
		308		FPV VP39	1.4c-135 1.7c-96	79/144 125/267	34	(Senkevich et al., 1997) (Binns et al., 1988)
088R	77075	185	21.3	RNA pol subunit rpo22				
J4R L4R	77632	185 185		VAC	1.2e-125	185/185	100	(Broyles and Moss, 1986) (Goebel et al., 1990)
		185		VAR-BSH myxoma	7.9e-125 1.5e-86	182/185 124/185	98	(Shchelkunov et al., 1995)
MC077R		187 186		MCV subtype I	1.9e-76	73/132	55	(Jackson and Bults, 1990) (Senkevich et al., 1997)
089L	78101	133	15.2		2.1e-73	72/135	53	(Binns et al., 1988)
JSL	77700	133	15.2	15.2k protein VAC	2.4e-95	122/122		(Plucienniczak et al., 1985)
L5L MC078L		133 134		VAR-I	2.4e-94	133/133 131/133	100 98	(Goebel et al., 1990) (Shehelkunov et al., 1995)
		137		MCV subtype I FPV	5.7c-45 1.4c-43	60/127 60/130	4/	(Senkevich et al., 1997)
		377 378		VAR-I A16L (BSH:A17L) VAC A16L	0.049	7/28	25	(Drillien et al., 1987) (Shehelkunov et al., 1995)
090R	78207		144.0		0.049	7/28	25	(Goebel et al., 1990)
J6R	82067	1286 1286	146.9	RNA pol subunit rpo147 VAC	0.0	1787/1707	00	(Broyles and Moss, 1986)
L6R MC079R		1286 1289		VAR	0.0	1283/1286	77	(Gnebel et al., 1990) (Shehelkunov et al., 1995)
		. 207		MCV subtype ( 100 matches to RNA pol (large	0.0 <3.7e-07	556/760		(Senkevich et al., 1997)
				subunit) family	- 45.76-07			
091L	82579	171	19.7	protein tyrosine/serine				(Rosel et al. 1994)
HIL	82064	171		phosphotase VAC	200 110	170/10:		(Rosel et al., 1986) (Guan et al., 1991)
111		171		VAR	2.0c-117 1.1c-114	170/171 166/171	99 97	(Goebel et al., 1990) (Shchelkunov et al., 1995)
		172		racoonpox myxoma virus	6.0c-111	157/171	91	847452
MC082L		173 169		rabbit fibroma virus	1.8c-77	83/138 46/80	60 57	(Mossman et al., 1995a) (Mossman et al., 1995a)
				MCV subtype I protein phosphatase family	1.4e-65 >2.8e-05	60/114	52	(Senkevich et al., 1997)
	<del></del>							

ORF	START		kDa'	name / (putative) function / homologies	BLAST expect	BLAST <sup>c</sup> AA id	HSS <sup>1</sup> (%)	references
left te	82593	regio 189	n: 21.5	21.5k protein				(Perel at 1996)
092K H2R	82393	189	21.5	21.5k protein VAC	5.2c-134	188/189	99	(Rosel et al., 1986)
I2R		189		VAR	1.4c-133	188/189	99	(Goebel et al., 1990) (Shcheikunov et al., 1995)
MC083R		191		MCV subtype 1	1.4e-71	95/181	52	(Senkevich et al., 1997)
		142		myxoma	1.3e-65	93/142	65	(Jackson and Bulis, 1990)
093L	84139	324	37.5	immunodominant env				(Posel at al. 1096)
0,52	83165	324	37.2	protein p35; IMV				(Rosel et al., 1986) (Chertov et al., 1991)
			_	membrane-associated				(Takahashi et al., 1994)
H3L		324	` .	VAC	3.3e-231	322/324	99	(Goebel et al., 1990)
I3L		325		VAR-BSH	1.7e-225	311/320	97	(Shchelkunov et al., 1995)
MC084L		298		MCV subtype 1	1.1c-36	38/117	32	(Senkevich es al., 1996)
094L	86527	795	93.6	RAP 94 (RNA-pol assoc.				(Ahn and Moss, 1992b)
	84140			transer. spec. factor)				(Kane and Shuman, 1992)
H4L		795		VAC VAR	0.0	791/795	99	(Goebel et al., 1990)
I4L MC085L		795 791		MCV subtype 1	0.0 0.0	780/795	98 59	(Shchelkunov et al., 1995)
MICOOJE		804		Orf virus	0.0	327/546 96/131	73	(Senkevich et al., 1996) (Fleming et al., 1993)
		484		FPV LIL protein	2.4c-181		51	2209386A
095R	86713 87324	203	22.3	late transcription factor VLTF-4				(Kovacs and Moss, 1996)
H5R		203		VAC	1.8e-128	202/203	99	(Rosel et al., 1986) (Goebel et al., 1990)
15R		221		VAR	5.1c-102	91/97	93	(Shchelkunov et al., 1995)
		227		orf virus F3R	· 3.1e-14	29/69	42 .	(Fleming et al., 1993)
		220		MCV subtype I	3.1c-09	28/64	43	(Senkevich et al., 1997)
•		705		nucleolin Xenopus	0.00041	18/57	31	(Messmer and Dreyer, 1993)
				31 matches to glu/asp rich proteins	E<0.52			
				p. 0.0.03				
096R	87325	314	36.7	DNA topoisomerase I				(Shuman and Moss, 1987)
U.C.D	88269	214		· VAC	0.0	214/214		(Rosel et al., 1986)
<i>H6R</i> 16R		314 314		VAC VAR-BSH	0.0 9.5e-220	314/314 312/314	100 99	(Goebel et al., 1990)
		314		shope fibroma virus	8.5e-141	119/170	70	(Shchelkunov et al., 1995) (Upton et al., 1990b)
		318		orf virus	5.2e-128	82/138	59	(Fleming et al., 1993)
MC087R		323		MCV subtype I	1.6c-121	111/202	54	(Senkevich et al., 1997)
		316		FPV L3R	2.9e-113	159/303	52	(Zantinge et al., 1996)
				21 matches to topoisomerase				
				family				
097R	88306	146	17.0	17.0k protein				(Rosel et al., 1986)
H7R	88746	146	•	VAC	2.1e-98	144/146	98	(Goebel et al., 1990)
17R		146		VAR	6.70-96	141/146	96	(Shehelkunov et al., 1995)
MC088R		143		MCV subtype I	4.3c-30	45/115	39	(Senkevich et al., 1997)
098R	88790	844	96.8	mRNA capping enzyme,				(Morgan et al., 1984)
	91324			large subunit				(Niles et al., 1986)
DIR		844		VAC	0.0	842/844	99	(Goebel et al., 1990)
FIR		844		VAR-BSH	0.0	830/844	98	(Shchelkunov et al., 1995)
MCO90R		950 836		MCV subtype t shope fibroma virus	0.0 0.0	322/64 243/305	64 79	(Senkevich et al., 1997) (Upton et al., 1991b)
		868		ASV NP868R	0.0033	17/55	30	(Pena et al., 1993)
				•			-	,,,
099L	91723 91283	146	16.9	structural protein -				(Niles et al., 1986)
D2L	71205	146		VAC	5.9c-98	146/146	100	(Dyster and Niles, 1991) (Goebel et al., 1990)
F2L		146		VAR (BSH: F3L)	1.5e-97	145/146	99	(Shchelkunov et al., 1995)
		143		Rabbit fibroma virus	2.0€-27	13/33	39	(Upton et al., 1991b)
MC091L		170		MCV subtype 1	1.1c-20	19/41	46	(Senkevich et al., 1996)
100 R	91716	233	27.6	27k structural protein	•			(Dyster and Niles, 1991)
D3R	92417	237		VAC	3.8-167	136/142	95	(Goebel et al., 1990)
F2R		237		VAR I:F3R	1.5e-162	131/142	92	(Shchelkunov et al., 1995)
		241		shope fibroma virus	9.3e-20	27/100	27	(Upton et al., 1991b)
MC092R		268 206		MCV subtype I rabbit fibroma virus C3	3.5e-18 1.6e-09	16/39 26/96	4 I 27	(Senkevich et al., 1997) (Strayer et al., 1991)
		200		TABOR TIBIORIA VITAS CS	1.00-03	20/70	21	(Strayer er at., 1991)
101R	92417	218	25.1	uracil DNA glycosylase				(Upton et al., 1993)
D4R	93073	218		VAC	1.4e-157	217/218	99	(Goebel et al., 1990)
F4R		218 218	•	VAR-BSH shope fibroma virus	5.1c-157 1.5c-117	216/218 151/218	99 69	(Shcheikunov et al., 1995) (Upton et al., 1993)
MC093R		226		MCV subtype 1	8.4c-91	65/113	57	(Senkevich et al., 1997)
		218		FPV FPD4	3.1e-88	116/216	53	(Tartaglia et al., 1990)
		297		uracil DNA glycosylase UL2	0.019	8/14	57	L34064
				gallid herpesvirus 1				
102R	93105	785	90.4	90.4k ATP/GTP binding				(Niles et al., 1986)
	95462			protein				(Shchelkunov et al., 1993c)
D5R		785		VAC	0.0	780/785	99	(Goebel et al., 1990)
FSR		785		VAR	0.0	774/785	98	(Shchelkunov et al., 1995)
	•	786 791		shope fibroma C5 MCV subtype (	0.0 0.0	283/450 184/334	62	(Strayer et al., 1991)
		791		FPV virus FPD5	0.0	184/334 170/345	55 49	(Senkevich et al., 1997) (Tartaglia et al., 1990)
MC094R		942		C29E6.4 C. clegans	0.72	16/56	28	(Wilson et al., 1994)
1025	05502	625	77.0	•				
103R	95503 97416	637	73.9	early transcription factor VETF-1	•			(Broyles and Fesler, 1990) (Gershon and Moss, 1990)
								(-cramon and moss, 1330)

-38-GENOMIC SEQUENCE OF THE MVA STRAIN

	ORF	STA STO	RT A	kD:	(20000170)	BLAST		HSS	
For	left			ion:	function / homologies	expec	t AA id	(%	)
MCD99R					VAC	0.0	635/637	90	(Carlot
MCV subype   605	F6R					0.0			(Shehelkunov et al. 1996)
Company   Comp	MC09:	5R			MCV subtune t				(Strayer et al., 1991)
Carugha et al., 1990			605	;					(Senkevich et al., 1997)
104K   97443   61   17.9   RA polymerase subunit (ever vitus)   1.5-0.5   13/38   34   (Yanz ad Moyer, 1991)			648		Charistaneuss biomic cau				(Tartaglia et al. 1990)
104R					Amsacta moorei EPV				(Yuen et al., 1991)
1048			706		African swine fever virus				(Yanez et al. 1992)
## State	104R	97443	161	17.9	RNA polymerase				
## MC097R	Dan	97928			subunit epo18		-		(Ahn et al., 1990b)
MCO97R   163									(O00001 et al., 1000)
MCD97R			163						(Suchelkunov et al. 100c)
105L   98805   97891   98805   97891	MC097	R			MCV subtype 1	4.0c-70			(Senkevich et al 1907)
PRE					FFV D/	5.4c-66	95/160	59	(Binns et al., 1990)
DBL	105L		304	35.4	virion transmembrane	,			(Niles and Seto 1000)
DRL		97891							(Niles et al., 1986)
10   10   10   10   10   10   10   10						2.3c-212	297/304 0	7	(Maa et al., 1990)
106	FaL					2.5c-209	291/304 9:	5	(Shchelkunov et al. 1995)
106 R									X97857
106R   9847   213   25.0   25k muft-like protein   23k-304   93					Monkeypox virus				
10			304				285/304 93	3	
99488  99488  213  VAC  1.6c-146  212/13  99 (Gondin, 1993) (Koonin, 1993) (Gosbel et al., 1996) (Gosbel et al., 1997) (Konin, 1993) (Kliss et al., 1986) (Gosbel et al., 1997) (Gosbel et al., 1997) (Konin, 1993) (Kliss et al., 1986) (Gosbel et al., 1997) (Gosbel et al., 1997) (Konin, 1993) (Kliss et al., 1986) (Konin, 1993) (Kliss et al., 1987) (Konin, 1993) (Kliss et al., 1987) (Konin, 1993) (Kliss et al., 1986) (Kliss	1000	000.0			·	>4.96-13			
D9R	106K		213	25.0	25k mutT-like protein				(Koonin, 1993)
MC098R   218		,,,,,,			VAC	I 6c-146	212/213	90	(Niles et al., 1986)
MCO99R 212 MCV subtype 1	F9R					5.3e-145			(Shchelkunov et al. 1995)
MCO99R	MC0981	3			MCV subtype I				(Strayer et al., 1991)
MCV subtype     0.0041   13/31   41   (Senkevich et al., 1997)	MCGGG				FPV D9				(Senkevich et al., 1997)
2215	MCO99F	•			MCV subtype 1 .			41	(Senkevich et al., 1997)
VAC DIOR   0.23   11/26   42   (Goebel et al., 1990)   (Robert et al., 1990)			225		FPV DIO				(Shchelkunov et al., 1995)
107			248		VAC DIOR				(Gocbel et al., 1990)
DIOR	107R		248	28.9	29k mutT-like protein				
FIOR 248 VAR-I	DIOR	100231	210						
MC099R   229									(Goebel et al., 1990)
MCV subtype     1.4e-54   44/100   44   (Senkevich et al. 1997)	MCOOOD				shope fibroma D10				(Straver et al., 1995)
218   shope fibroms D9   1.9e-06   195-4   35   31102   44   44   45   45   45   45   45   4	MICUSSK	•							(Senkevich et al., 1997)
MCV subtype   MCO98R   0.13   12/21   57   (Senkevich et al., 1997)   (Senkevich et al., 1996)   (Se			218		shope fibroma D9				(Binns et al., 1990)
108L   102127   631   72.4   nucleoside triphosphute phosphotydrolase 1,					MCV subtype i MC098R	0.13	12/21	57	(Senkevich et al., 1997)
169			213	•					D90899
108L   102127   631   72.4						0.24			(Shehelkunov et al. 1995)
100232			103		mutator M. Jannaschii	0.39	13/25	52	(Bult et al., 1996)
DILL   631	108L		631	72.4					(Brayles and Moss 1997)
D71L   631		100232			phosphohydrolase I, DNA helicase				(Rodriguez et al., 1986)
MC100R 631 VAR 0.0 626/631 99 (Shchelkunov et al., 1995)  MC100R 634 MCV subtype 1 7.3c-286 392/627 62 (Senkevich et al., 1996)  MC100R 637 FPV protein 5 2.8e-275 214/357 59 542251  MC100R 638 MCV subtype 1 7.3c-286 392/627 62 (Senkevich et al., 1996)  MC100R 634 MCV subtype 1 7.3c-286 392/627 62 (Senkevich et al., 1996)  MC100R 637 FPV protein 5 2.8e-275 214/357 59 542251  MC100R 638 MCV subtype 1 7.3c-286 392/627 62 (Senkevich et al., 1996)  MC100R 634 MCV subtype 1 7.3c-286 392/627 62 (Senkevich et al., 1996)  MC100R 634 MCV subtype 1 7.3c-286 392/627 62 (Senkevich et al., 1996)  MC100R 634 MCV subtype 1 7.3c-286 392/627 62 (Senkevich et al., 1996)  MC100R 634 MCV subtype 1 7.3c-286 392/627 62 (Senkevich et al., 1993)  MC100R 637 FPV protein 6 7 5.4e-286 394/287 99 (Shchelkunov et al., 1993)  MC101L 103025 287 Swinepox virus 4.1c-160 220/287 76 (Massung et al., 1993)  MC101L 104711 551 61.9 rifampicin resistance gene, IMV protein  MC102L 547 MCV subtype 1 5.4e-286 171/279 61 (Senkevich et al., 1996)  MC102L 547 MCV subtype 1 5.4e-286 387/551 99 (Massung et al., 1995)  MC102L 547 MCV subtype 1 5.4e-286 298/494 60 (Massung et al., 1995)  MC102L 547 MCV subtype 1 5.4e-286 298/494 60 (Massung et al., 1995)  MC102L 547 MCV subtype 1 5.4e-286 298/494 60 (Massung et al., 1995)  MC102L 547 MCV subtype 1 5.4e-286 298/494 60 (Massung et al., 1995)  MC102L 547 MCV subtype 1 5.4e-286 298/494 60 (Massung et al., 1995)  MC102L 547 MCV subtype 1 5.4e-286 298/494 60 (Massung et al., 1995)  MC102L 547 MCV subtype 1 5.4e-286 298/494 60 (Senkevich et al., 1996)  MC102L 548 MCV subtype 1 5.4e-286 298/494 60 (Senkevich et al., 1996)  MC102L 548 MCV subtype 1 5.4e-286 298/494 60 (Senkevich et al., 1996)  MC102L 548 MCV subtype 1 5.4e-286 298/494 60 (Senkevich et al., 1996)  MC102L 548 MCV subtype 1 5.4e-286 298/494 60 (Senkevich et al., 1996)						0.0	629/631 99		(Koonin and Senkevich, 1992)
Section   Sect						0.0	626/631 99		(Shchelkunov et al., 1995)
Sample   S									Senkevich et al., 1996)
Charistoneura biennis EPV   1.1e-136 81/158 51   Cyten et al., 1991)					Rabbit fibroma C14 protein	1.8c-176			
Swinepox virus   1.2e-34   60/89   67   (Massung et al., 1993)   1085   RAD26 (yeast)   5.1e-05   16/45   35   (Huang et al., 1993)   (Huang et al., 1994)   (Okabe et al., 1995)   (Okabe et al., 1995)   (Okabe et al., 1996)   (Okabe et al., 1995)   (Okabe et al., 1996)   (Okabe et al., 1995)   (Okabe et al., 1996)   (Okabe et al., 1995)   (Okabe et al., 1996)   (								9	Hall and Moyer, 1991)
1098					Swinepox virus				Yuen et al., 1991) Massung et al. 1993)
HS transcription activator   0.00093   10/22   45   (Okabe et al., 1994)   (Okabe et al., 1992)							26/89 29	(	Baylis et al., 1993)
NTPase family >5.1c-5  NTRA capping enzyme, transcription initiation factor VITF  NAC   2.0c-198   285/287   99   (Weinrich and Hruby, 1986) (Vos et al., 1991)   (Vos et al., 1990)   (Vos et al., 1991)   (Vos et al., 1995)   (Vos et al., 1995)   (Nossung et al., 1995)   (Nossung et al., 1995)   (Nossung et al., 1996)   (Nossung et al., 1985)   (Nose et al., 1996)   (Nose et al., 1996)   (Nossung et al., 1995)   (Nose et al., 1996)   (No					HS transcription activator				
102162					NTPase family		,,,	. `	Chabe 21 at., 1992)
102162   transcription initiation   (Weinrich and Hruby, 1986)   (Weinri	109L		287	33.3	mRNA capping enzyme.		•	٠.	Niles et et 1000
D12L   287		102162			transcription initiation			(	Weinrich and Heihy 1996)
N2L 287 VAR 9.8e-198 284/287 99 (Goebel et al., 1990) 287 Swinepox virus 4.1e-160 220/287 76 (Massung et al., 1993) 289 FPV protein 6 3.4e-113 114/215 53 (Senkevich et al., 1996) 3.4e-113 114/215 53 (Senkevich et al., 1995) 3.4e-113 114/215 53 (Senkevich et al., 1996) 3.4e-113 114/215 53 (S			287			7 Oc-100	285/207 ""	(	vos et al., 1991)
MCIOIL 287 Swinepox virus 4.1e-160 220/287 76 (Massung et al., 1993) 289 FPV protein 6 3.4e-113 114/215 53 S42252  III	N2L				VAR	9.8c-198	284/287 99	(	Shehelkunov et al. 1990)
289 FPV protein 6 3.6e-120 171/279 61 (Senkevich et al., 1996)  110L 104711 551 61.9 rifampicin resistance gene, IMV protein  D13L 551 VAC 0.0 551/551 100 (Goebel et al., 1998)  N3L 551 VAR 0.0 547/551 99  MC102L 547 MCV subtype 1 5.4e-248 298/494 60 (Senkevich et al., 1993)  MC102L 547 MCV subtype 1 5.4e-248 298/494 60 (Senkevich et al., 1993)  S52 FPV protein 7 6.6e-223 182/305 59 S42253	MCIOIL				MCV subsyne !	4.1c-160	220/287 76	- (	Massung <i>et al.</i> , 1993)
101	•							. (	Senkevich et al., 1996)
103056	110L	[047]]	551	61 9	rifamniain				
DI3L   S51   VAC   0.0   S51/551   100   (Weinrich and Hriby, 1986)     N3L   S51   VAR   0.0   S47/551   99   (Shehelkunov et al., 1995)     N51   Swinepox virus   4.5e-286   357/506   70   (Massung et al., 1993)     S52   FPV protein 7   6.6e-223   182/305   59   S42253     S84   Helightic regularies   FPV   S86   Helightic r				J,				Č	Tartaglia and Paoletti, 1985)
Solution					VAC			6	weinrich and Hruby, 1986) Goebel <i>et al.</i> , 1990)
MCI02L 547 MCV subtype 1 5.4e-248 298/494 60 (Massung et al., 1993) 552 FPV protein 7 6.6e-223 182/305 59 S42253								(:	Shchelkunov et al., 1995)
532 Prv protein 7 6.6e-223 182/305 59 \$42253	MC102L		547		MCV subtype 1	5.4e-248	298/494 60	(1	viassung et al., 1993) Senkevich et al. 1996)
Osborne et al., 1996)						6.6c-223	182/305 59	S	42253
	·					50-51	77/10/ 30	((	Usborne er al., 1996)

ORF'	STAR'S STOP	' AA	kDae	name / (putative) function / homologies*	BLAST <sup>d</sup> expect	BLAST	HSS <sup>(</sup>	references
	erminal	regio						
111L	105187	150	16.9	late gene trans-activator, VLTF-2				(Weinrich and Hruby, 1986)
AIL		150		VAC	6.8e-103	149/150 9		(Keck et al., 1993) (Goebel et al., 1990)
AIL MC103L		150		VAR	6.8e-103	149/150 9		(Shchelkunov et al., 1995)
WC 103L		169 154		MCV subtype I FPV protein 8	6.3e-54 2.8e-50	74/147 5 50/87 5		(Senkevich et al., 1996) \$42254
112L	105882	224	26.3					
AZL	105208		20	VAC				(Weinrich and Hruby, 1986) (Passarelli et al., 1996)
A2L		224		VAR	1.3e-158	224/224 1		(Goebel et al., 1990)
MC104L		224 228		MCV subtype 1 orf virus	1.3e-158	224/224 1		(Shchelkunov et al., 1995)
		606		5 <b>V</b> 1.05	6.4e-127 6.8e-30	172/222 7° 43/66 6.		(Senkevich et al., 1996) (Mercer et al., 1995)
113L	106109	76	8.9	8.9k protein				
	105879	76		VAC-WR	1.6e-47	73/76 9	6	(Weinrich and Hruby, 1986)
A3L MC105L		76 70		VAR-BSH (I:A2.5L)	2.1e-47	71/76 9:		(Shchelkunov et al., 1995)
				MCV subtype 1	9.8c-12	26/63 4	'	(Senkevich et al., 1996)
114L <i>A3L</i>	108058	644 644	72.6	major core protein P4b VAC	0.0	6431644 0		(Rosel and Moss, 1985)
A4L	100121	644		VAR-BSH (I:A3L)	0.0 0.0	643/644 9		(Goebel et al., 1990) (Shchelkunov et al., 1995)
MC106L		675		MCV subtype 1	8.9e-272	227/357 6	3	(Senkevich et al., 1996)
		657		FPV Major core protein P4b	9.1c-220	169/299 5	6,	(Binns et al., 1989)
115L	108929	272	29.9	months and appointed fort				(Demkowicz et al., 1992)
A4L	108111	281		protein VAC	1,1e-145	180/187 96	6	(Cudmore et al., 1996)
A5L		271		VAR-BSH (I: A4L)	1.1e-112	165/178 9:		(Goebel et al., 1990) (Shchelkunov et al., 1995)
		268 5179		Thermoproteus phage 1 human mucin	1.9e-09 4.5e-07	38/127 25		(Neumann and Zillig, 1990)
		3.,,		many matches to Pro-rich	4.30-07	34/139 2	4	(Gum et al., 1994)
				proteins				
116R	108967	164	19.0	RNA pol subunit rpo19				(Ahn et al., 1992)
A5R A5R	109461	164		VAC	5.8c-110	164/164	100	(Goebel et al., 1990)
MC108R		164 165		VAR-I (BSH:A6R) MCV subtype 1	7.0e-109 3.3e-51	162/164 82/151	98 53	(Shehelkunov et al., 1995)
		167		FPV	3.3e-51	72/161	44	(Senkevich et al., 1997) (Kumar and Boyle, 1990)
				54 matches/glu-rich proteins	<0.51			, ,
· 117L	110576	372	43.1	43.1k protein			_	
<i>A6L</i> A7L	109458	372 372		VAC VAR-BSH (I: A6L)	1.2e-248 1.1e-244	371/372 99 364/372 91		(Goebel et al., 1990)
MC109L		461		MCV subtype !	4.0c-99	132/367 35		(Shchelkunov et al., 1995) (Senkevich et al., 1996)
		339		FPV ORF 2 protein	1.9e-95	111/279 39		B60013
118L	112732		82.3	VETF 82k subunit				(Gershon and Moss, 1990)
<i>A7L</i> A8L	110600	.710 710		VAC VAR-BSH (I: A7L)	0.0	708/710 99	9	(Goebel et al., 1990)
WCITOL		707		MCV subtype !	0.0	700/710 98		(Shchelkunov et al., 1995) (Scnkevich et al., 1996)
119R	112786	288	33.6	33.6k protein				
A 8 R	113652	288		VAC	5.3e-198	287/288	99	(Van Meir and Wittek, 1988) (Goebel et al., 1990)
A8R MCIIIR		288 435		VAR-I (BSH:A9R)	3.1e-195	284/288	98	(Shchelkunov et al., 1995)
				MCV subtype 1	4.4e-94	100/169	59	(Senkevich et al., 1997)
120L A10L	113929 113645	94	10.5	10.5k protein		70/70 31		(Van Meir and Wittek, 1988)
A9L	.1.7043	95 99		VAR-BSH (I: A9L) VAC	9.0e-59 9.4e-55	78/79 98 82/91 90	_	(Shchelkunov et al., 1995) (Gochel et al., 1990)
MC112L		128		MCV subtype I	1.0e-29	47/71 66	5	(Senkevich et al., 1996)
		69		orf virus	3.0e-16	27/45 60	ט	(Mercer et al., 1995)
121L	116605	168	102.2	major core protein P4s				(Van Meir and Wittek, 1988)
AIOL	113930	891		VAC	0.0	883/891 99		(Vanslyke et al., 1991) (Goebel et al., 1990)
AIIL		892		VAR-BSH (I: A10L)	0.0	442/463 95	5	(Shchelkunov et al., 1995)
MCH3L		889		MCV subtype 1	5.8e-289	99/177 55	5	(Senkevich et al., 1996)
122R	116620	318	36.1	36.1k protein				(Goebel et al., 1990)
AIIR AIIR	117576	318		VAC VAR-I (BSH: AI2R)	3.5e-212 2.7e-154	318/318 242/277	100 87	(Gocbel et al., 1990)
MC114R		304		MCV subtype !	2.9e-98	92/154	59	(Shchelkunov et al., 1995) (Senkevich et al., 1997)
		148		FPV 4u gene	1.9e-13	18/32	56	A20158
123L	118141	187	20.0	virion protein			•	(Takahashi <i>et ul.</i> , 1994)
A12L A13L	117578	192 189		VAC VAR-BSH (I: A12L)	4.8e-127 5.9e-64	127/128 99	•	(Goebel et al., 1990)
MCIISL		178		MCV subtype 1	5.9e-04 5.9e-37	39/83 46		(Shchelkunov et al., 1995) (Senkevich et al., 1996)
124L	118377	70	7.6	structural protein				
	118165			IMV membrane protein			,	(Takahashi <i>er al.</i> , 1994) (Jensen <i>er al.</i> , 1996)
A/3L		70 68		p 8 VAC	2.4c-42	66/69 95	5	(Goebel <i>et al.</i> , 1990)
A14L				VAC VAR-BSH (I: AI3L)	4.1c-20	37/64 57	'	(Shchelkunov et al., 1995)
125L	118757	90	10.0	structural protein				(Takabashi wast 1994)
	118485	,,	10.0	IMV membrane protein				(Takahashi <i>et al.</i> , 1994) (Jensen <i>et al.</i> , 1996)
				p 1 6				

-40-GENOMIC SEQUENCE OF THE MVA STRAIN

ORF*		Т АЛ	kDa'	mand , (pullitie)	BLAST	BLAST	* HSS	references
left	STOP terminal	regi	оп:	function / homologles*	expect	AA Id	(%	
A14L A15L MC118	iL	90 90 94 125		VAC VAR-BSH (I: A14L) MCV subtype 1 human interferon inducible protein	5.3e-62 5.3e-61 7.3e-22 0.23	90/90 88/90 31/72 15/49	100 97 43 30	(Goebel et al., 1990) (Shcheikunov et al., 1995) (Senkevich et al., 1996) (Deblandre et al., 1995)
126L <i>A15L</i> A16L MC120	11920 11892		11.0	IIk protein VAC VAR-BSH (I:AISL) MCV subiype !	4.1c-63 1.0e-61 6.7c-08	94/94 92/94 17/51	100 97 33	(Goebel et al., 1990) (Shehelkunov et al., 1995) (Senkevich et al., 1996)
127L A16L A17L MC121	119193	377 378 377 364	43.4	35k myristylprotein VAC VAR-BSH (I:A16L) MCV subtype i	6.3e-288 1.5c-283 6.5e-110	368/377	97	(Martin et al., 1997) (Gocbel et al., 1990) (Shchełkunov et al., 1995) (Senkevich et al., 1996)
128L	120940 120329		23.0	IMV membrane protein morphogenesis factor				(Krijnse-Locker et al., 1996) (Rodriguez et al., 1995)
A17L A18L MC122	د	203 203 179		VAC VAR-BSH (l:A17L) MCV subtype l	1.0e-141 1.0e-141 1.4e-47			(Wolfe et al., 1996) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1996)
129 R A ! 8 R A ! 8 R MC ! 23 I	12095 <i>5</i> 122436		56.8	DNA helicase DNA dependent ATPose VAC VAR-1 (BSH:A19R) MCV subtype 1 Bacteriophage T5 D10 helicase-like protein	0.0 0.0 5.3e-167 0.0066	488/493 478/493 203/403 13/36	96	(Koonin and Senkevich, 1992) (Bayliss and Condit, 1995) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1997) P11107
130 L <i>A19 L</i> A19 L MC124 L	122650 122417		8.3	8.3kb protein VAC VAR-I (BSH: A20L) MCV subtype I HS RIZ, zink finger protein	2.9c-50 1.2c-34 1.5c-13 0.0060	77/77 54/64 14/37 7/16	100 84 37 43	(Goebel et al., 1990) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1996) (Buyse et al., 1995)
131 <b>L</b> A21L A22L MC125L	123004 122651	117 117 117 114	13.6	13.6k protein VAC VAR-BSH (I: A20L) MCV subtype 1	5.3e-83 7.2e-82 2.8e-28	117/117 115/117 23/41		(Goebel et al., 1990) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1996)
132R A20R A21R MC126R	123003 124283	426 426 426 476 1118	49.1	49.1k protein 'VAC 'VAR MCV subtype 1 Pichia klyveri DNA pol	7.6e-298 1.6e-294 3.2e-95 0.069	423/426 418/426 34/131 12/54	99 98 25 22	(Goebel et al., 1990) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1997) Y11606
133R A22R A22R MC127R	124776	187 187 176 282	21.9	21.9k protein VAR-I (BSH:A23R) VAC MCV subtype 1	1.1e-126 1.2e-122 5.8e-43	182/187 174/176 35/85	97 98 41	(Goebel et al., 1990) (Shchelkunov et al., 1995) (Goebel et al., 1990) (Senkevich et al., 1997)
134 R <i>A23 R</i> A23 R MC128 R		382 382 382 383	44.6	44.6k protein VAC VARI (BSH:A24R) MCV subtype !	4.2e-269 1.7e-265 3.5e-136	382/382 377/382 83/143	100 98 58	(Goebel et al., 1990) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1997)
135R A24R A25R MC129R	125966 129436	1155 1164 1164 1164 1165 1162	132.4	VAC CPX rpo132 VAR-BSH (I:A24R) MCV subtype I orf virus 101 matches to RNA pol beta subunit family	0.0 0.0 0.0 0.0 0.0 0.0 <0.036	794/796 794/795 789/795 441/565 166/258	99 99 99 78 64	(Hooda-Dhingra et al., 1990) (Amegadzie et al., 1991b) (Goebei et al., 1990) (Patel and Pickup, 1989) (Shehelkunov et al., 1995) (Senkevich et al., 1997) U33419
	erminal			`				
136L A25L	129638 129441	65 65 1284	7.5	150k CPX-ATI (f) VAC Cowpox (CPX-ATI)	1.3e-41 3.2e-15		98 93	(Funahashi et al., 1988) (Goebel et al., 1990) (Funahashi et al., 1988)
137L A30L A26L MC131L MC133L MC130L	130916 130224	230 498 322 513 546 451 702 726	27.1	27.1k protein (f) VAR-BSH (l: A29L) VAC (AT1 flanking protein) MCV subtype I MCV subtype I MCV subtype I VAR-I A28L (BSH:A29L) Camelpox	3.1c-158 5.6c-142 2.1c-12 4.2c-11 2.3c-10 0.0021 0.051	12/40 14/40 12/37	98 32 30 35 32	(Amegadzie et al., 1991a) (Shchelkunov et al., 1995) (Goebel et al., 1990) (Senkevich et al., 1996) (Senkevich et al., 1996) (Senkevich et al., 1996) (Senkevich et al., 1996) (Shchelkunov et al., 1995) (Meyer and Rziha, 1993)
138L A27L A31L	131298 130966	110 110 110 117 110	12.5	14k membrane protein EEV protein fusion protein VAC VAR-BSH (I: /A30L) Camelpox virus Cowpox virus	3.3e-70 1.1e-69 1.5e-69 1.6e-69	108/110 9 107/110 9 106/110 9 107/110 9	98 97 96	(Rodriguez and Esteban, 1987) (Rodriguez and Smith, 1990) (Gong et al., 1990) (Goubet et al., 1990) (Shchelkunov et al., 1995) (Meyer et al., 1994) (Meyer et al., 1994)

ORF.		RT AA	kDs	' name / (putative)	BLAST	" BLAST	· us	(S)(S)
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		110	)	Ectromelia virus	6.7e-68	105/110	9 9 5	(Manual L. 1994)
		110 89	)	Monkeypox virus Orf virus	8.3e-67	103/110	93	(Meyer et al., 1994) (Meyer et al., 1994)
	_	188		Myxoma virus	4.8c-15 2.5c-12		38 54	(Naase et al., 1991) (Jackson et al., 1996)
MCI33	L	546 148		MCV subtype 1 Capripox virus HM2 protei	1.5e-11	26/58	44	(Senkevich et al., 1996)
MC131	L	513		MCV subtype 1	n 2.6e-10 1.5e-05	21/42 18/58	50 31	(Gershon et al., 1989) (Senkevich et al., 1996)
139L	13173			16.3k protein				(Amegadzie et al., 1991a)
A28L A31.5L	13129	9 146 146		VAC VAR-BSH (I: A31L)	1.7c-10:			(Goebel et al., 1990)
		140		Myxoma virus	2.9e-10 1.3e-55	0 141/146 30/52	96 57	(Shchelkunov et al., 1995) (Jackson et al., 1996)
MC1341	L	140 141		Capripox virus HM3 protei MCV subtype I	n 3.3e-55 1.0e-53	30/49	61	(Gershon et al., 1989)
		143		Amsacta moorei poxvirus	2.0e-14	31/52 16/36	59 44	(Senkevich et al., 1996) (Hall and Moyer, 1991)
140L <i>A29L</i>	13265			RNA pol subunit rpo35				(Amegadzie et al., 1991a)
A32L	13174	305		VAC VAR-BSH	3.6e-215 7.5e-211			(Goebel et al., 1990)
MC1351	٠.	303 126		MCV subtype I	7.0c-98	51/103	49	(Shchelkunov et al., 1995) (Senkevich et al., 1996)
				Capripox virus	2.2e-54	46/61	75	(Gershon et al., 1989)
141L <i>A30L</i>	13285		8.7	8.7k protein VAC				(Amegadzie et al., 1991a)
A33L		77		VAR	5.5c-48 5.5c-48	77/77 · . 77/77	001	(Goebel et al., 1990)
MC136L	•	67		MCV subtype 1	9.2e-16	18/40	45	(Shchelkunov et al., 1995) (Senkevich et al., 1996)
142R <i>A31R</i>	133013		14.4	14.4k protein				(Smith et al., 1991)
A34R	133390	124		VAC VAR	2.0c-84 1.6c-79	118/124	95	(Goebel et al., 1990)
MC138R	t	117		MCV subtype 1	6.2e-24	111/114 39/98	·97 39	(Shchelkunov et al., 1995) (Senkevich et al., 1997)
143L	134 169 133360		30.8	30.8k protein				(Smith et al., 1991)
A32L	133300	300		ATP/GTP binding motif	A 6.4c-190	268/269	99	(Koonin et al., 1993)
A35L MC140L		270 249		VAR	1.6e-186	263/269		(Goebel et al., 1990) (Shchelkunov et al., 1995)
				MCV subtype I	7.6e-95	58/94	61	(Senkevich et al., 1996)
144 R A33R	134287 134844		20.6	EEV glycoprotein VAC	2.1.104			(Roper et al., 1996)
A36R		184		VAR	2.1c-124 1.8c-121	182/185	8e 1e	(Goebel et al., 1990) (Shehelkunov et al., 1995)
		185		Ectromelia	2.8e-113	165/185	89	(Roper et al., 1996)
145R	134868 135374	168	19.6	EEV glycoprotein virulence factor				(Duncan and Smith, 1992a)
A 34 R		168		actin microvilli induces				(McIntosh and Smith, 1996) (Wolffe et al., 1997)
A37R		168		VAR-I	1.2e-117 1.7e-117	165/168 164/168	98 97	(Goebel et al., 1990)
		167		FPV ORFs BamHI 2,8,11 hepatelectins homologs	tic <0.056	16/66	24	(Shchelkunov et al., 1995) (Tomley et al., 1988)
		199		HS early T-cell activation	0.0038	12/38	31	(Hamann et al., 1993)
MCI43R		159		antigen CD69 MCV subtype 1	0.080	-		
				17 matches to lectins	0.000	12/48	25	(Senkevich et al., 1997)
146R <i>A35R</i>	135418 135948	176 176	20.0	20.0k protein				(Smith et al., 1991)
A38R	.55,46	60		VAC VAR-I	1.4c-126 2.9c-37	176/176 57/60	100 95	(Goebel et al., 1990)
MC145R		233		MCV subtype 1	1.2e-07	18/55	32	(Shchelkunov et al., 1995) (Senkevich et al., 1997)
147R	136015 136641	208	23.8k	EEV membrane protein				(Parkinson and Smith, 1994)
A36R		221		VAC	2.8e-143	140/141	99	(Smith et al., 1991) (Goebel et al., 1990)
A39R		216		VAR 19 matches to asn/ser-rich	2.1c-89 <0.41	138/177	77	(Shcheikunov et al., 1995)
				proteins	<b>NO.41</b>			
	136705	263	29.8	29.8k protein	,			•
<i>A37R</i> A40R	137496	263 68		VAC VAR	6.8c-183	261/262	99	(Goebel et al., 1990)
	120600				4.9e-37	61/67	91	(Shchelkunov et al., 1995)
	138589 137756	277 277	31.5	31.5k protein VAC	9.3e-198	274/277 9	10	(Amegadzie et al., 1991a)
A41L		277 303		VAR	1.6c-187	259/277 9		(Goebel et al., 1990) (Shchelkunov et al., 1995)
		324		Rattus norvegicus CD47 MM integrin assoc. protein	3.9e-24 1.0e-21		16 16	(Nishiyama et al., 1997)
		323		human CD47 precursor	5.0e-19		12	(Lindberg et al., 1993) (Campbell et al., 1992)
	138606 138857	83	9.4	semaphorin-like protein				(Kolodkin et al., 1993)
439R		403		VAC	8.0c-46	73/76	96	(Goebel et al., 1990)
	139163	74 210	23.9	VAR-i semaphoria-like protein	8.6e-44	67/71	94	(Shehelkunov et al. 1995)
	139795	403		(12)				(Kolodkin et al., 1993)
A43R		139		VAC VAR (I:A44R)	3.0-147 1.8c-68	209/210 91/105	99 86	(Goebel et al., 1990)
-		653		semaphorin-like protein Alcelaphine herpesvirus	1.76-20	29/79	36	(Shchelkunov et al., 1995) (Ensser and Fleckenstein, 1995)
				37 matches to semaphorin				
				•				

GENOMIC SEQUENCE OF THE MVA STRAIN

ORF'	STAF	RT AA	kDa	(putative)	BLAST		HSS	S' references
left	termina		ion:	/collapsin gene family	expec	AA Id	(%	)
152 R <i>A40R</i> A45R	13982 14032		3 3 2	NK cell receptor homolo lectin-like protein VAC VAR-I (BSH: A43.5R) HS natural killer (NK) cell protein group 2-A, B HS type II membrane protein MM NK cell receptor HS CD 94	6.6c-97 9.6c-36 4.5c-11	54/59 20/74	97 91 27 44 44 37	(Shchelkunov et al., 1995) (Houchins et al., 1991) (Adamkiewicz et al., 1994)
153L <i>A41L</i> A44L	14102 14036	5 219 6 219 218 244 258		127 matches to lectins including NK cell surface proteins and snake venoms  25.1k protein VAC VAR-BSH (I:A46L) VAC B29R/C23L Rabbit fibroma virus T1	1.9e-158 1.4e-152 0.0076 0.057	3 218/219	99 95 22 26	(Smith et al., 1991) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Goebel et al., 1990)
154R A42R A47R	14119 14158			profilin-like protein VAC VAR-I (BSH:A45R) HS. profilin 10 matches profilin family	1.2e-87 1.4e-85 2.2e-23	85/87 82/87 19/45	97 94 42	(Upton et al., 1987) (Blasco et al., 1991) (Smith et al., 1991) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Kwiatkowski and Bruns, 1988)
155R A43R A48R 156R	141621 142193	194 195 51	22.1	class I membrane glycoprotein VAC VAR-I(BSH:A46R) HS leukocyte antigen	1.5e-137 1.9e-128 0.096	162/164 101/109 7/23	98 92 30	(Smith et al., 1991) (Duncan and Smith, 1992b) (Goebel et al., 1990) (Shehelkunov et al., 1995) X79517
130K	142437		8.8	8.8k protein VAC-WR Salf6R rabbit myosin heavy chain 144 matches to various asp/glu/lys-rich proteins	3.9e-45 0.00048	78/78 13/39	100 33	(Smith et al., 1991) (Smith et al., 1991) A02985
157 L A44 L A47 L MC152R	143577 142537		39.4	3ß-hydroxysteroid dehydrogenase (3ß-HSD) VAC VAR-BSH (I: A49L) MCV subtype I FPV matches to dihydroflavonol reductases, cholesterol dehydrogenases, UDP- galactose-4-epimerases	4.5e-249 1.1e-136 8.2e-104 3.1e-83 >2.8e-05	342/346 185/195 123/272 33/85	98 94 45 38	(Moore and Smith, 1992) (Blasco et al., 1991) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Senkevich et al., 1996) (Skinner et al., 1994) (Baker and Blasco, 1992)
158R A45R A51R	143624 143989	121 125 125	13.3	superoxide dismutase-like protein VAC VAR-I BSH A48R 117 matches with superoxide dismutase family	2.1e-82 1.1e-82 <0.027	94/96 93/96	97 96	(Blasco et al., 1991) (Smith et al., 1991) (Goebel et al., 1990) (Shehelkunov et al., 1995)
159 R A46R A52R	143979 144701	241 214 240	27.6	27.6k protein VAC VAR-i (BSH: A49R)	9.6e-167 5.6e-164	238/240 233/240	99 97	(Smith et al., 1991) (Goebel et al., 1990) (Shchelkunov et al., 1995)
160L JIL <i>A47L</i>	145465 144749	238 244 244	27.6	27.6k protein VAR VAC integrin lipid binding motif	5.1c-146 8.2c-135	114/127 121/127	89 95	(Goebel et al., 1990) (Shchelkunov et al., 1995) (Goebel et al., 1990) (Smith et al., 1991)
161R <i>A48R</i> J2R	145564 146178	204 204 205	23.2	thymidytate kinuse VAC VAR 16 matches to thymidylate kinase family	5.2c-140 1.1c-137 <0.49	204/204 161/165	100 97	(Smith et al., 1991) (Goebel et al., 1990) (Shchelkunov et al., 1995)
162R <i>A49R</i> J3R	146202 146690	162 162 162	18.8	18.8k protein VAC VAR	6.0e-106 2.4e-103	159/162 154/162	98 95	(Smith et al., 1991) (Goebel et al., 1990) (Shchelkunov et al., 1995)
163R <i>ASOR</i> J4R	146722	552 552 552 922 559 564	63.5	DNA liguse VAC VAR-I HS DNA ligase III shope fibroma ligase FPV ligase 31 matches mainly to DNA ligase family	0.0 0.0 2.1e-235 9.9e-213 3.0e-195 <0.029	547/552 537/552 102/165 95/200 101/170	99 97 61 47 59	(Kerr and Smith, 1989) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Wei et al., ) (Parks et al., 1994) (Skinner et al., 1994)
	149358	310 334 334		34.9k protein VAC VAR	l.5e-217 9.1e-208	267/274 251/274	97	(Antoine et al., 1996) (Goebel et al., 1990) (Shchelkunov et al., 1995)

ORF	STAR	T AA	h kDa <sup>c</sup>	name / (putative) function / homologies*	BLAST			references
left	terminal	regi	on:		- Apect	AA id	(%)	
				fusion of ASIRIASSR ORFs				(Antoine et al., 1996)
165R <i>A56R</i> J9R	1494 (6 150363		34.8	hemagglutinin VAC VAR-I(BSH:J7R) raccoonpox 124 matches to various proteins	1.8c-211 4.3c-178 1.5c-91 <0.34		99 76 71	(Shida, 1986) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Cavallaro and Esposito, 1992)
166R <i>A57R</i> JIOR	150659 150952		11.4	guanylate kinase (f) VAC VAR (BSH:J8R) MM guanylate kinase HS guanylate kinase 21 matches mainly to guanylate kinases	3.2e-62 2.2e-57 4.3e-24 2.8e-20 <0.20	94/97 88/97 39/91 35/91	96 90 42 38	(Smith et al., 1991) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Brady et al., 1996) (Brady et al., 1996)
167R	151103 152005		34.3	serine/threonine protein kinase	ı			(Howard and Smith, 1989) (Banham and Smith, 1992)
BIR BIR	-	300 300 283		VAC VAR-I VAC B12R IOO matches mainly to protein kinase family	7.1e-215 2.7e-210 4.9e-49 <0.00031	298/300 289/300 27/53	99 96 50	(Lin et al., 1992) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Goebel et al., 1990)
168R	152144		11.5	24.6k protein (f1)				•
169 R	152434	149 143	16.1	VAC histone H2A pea 24.6k protein (f2)	8.5e-38 0.59	54/60 16/50	90 32	(Goebel et al., 1990) P40281 (Goebel et al., 1990)
B2R	152720	219		VAC	5.7e-86	124/128	96	(Goebel et al., 1990)
170R <i>B3R</i>	152917 153456	179 124 167 92	20.9	20.9k protein (f) VAC VAC WR VAR-GAR H5R	8.2e-33 5.3e-45 3.4e-06	51/56 51/56 19/28	91 91 67	(Goebel et al., 1990) (Smith et al., 1991) U18339
171R	153683	177	21.4	65k ank-like protein				(Howard et al., 1991)
B4R B6R 172R	154216 154107 155336	558 558 409	47.7 ·	virulence factor (f1) VAC VAR-I (BSH:BSR) 65k ank-like protein	8.5e-107 1.7e-98	151/154 140/154	98 90	(Mossman et al., 1996) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Howard et al., 1991)
B4R B6R		558 558 483 1765 516 574 574 574 582 668 237 472 474 446 437 634	. 4	virulence factor ((2) VAC VAR-I (BSH:B5R) MYX M-T5 protein MM ankyrin 3 orf virus VAC B18R VAR-I B19R HS KIAA0379 CPX host range gene VAC WR hr gene VAC MIL CPX OIL VAR OIL CPX DIL VAC C9L 159 matches including ankyrin proteins	2.4e-283 2.3e-270 5.5e-10 9.7e-10 1.8e-09 3.3e-09 5.1e-09 1.7e-08, 2.8e-08 5.1e-07 8.7e-07 8.7e-07 8.7e-07	195/201 185/201 19/57 22/54 16/47 11/23 19/72 20/52 14/47 15/47 23/81 22/61 23/81 8/27	97 92 33 40 34 47 26 38 29 31 28 36 28 29	(Mossman et al., 1996) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Mossman et al., 1996) (Peters et al., 1995) U34774 (Goebel et al., 1990) (Shchelkunov et al., 1995) AB002377 (Spehner et al., 1988) (Kotwal and Moss, 1988a) (Goebel et al., 1990) (Safronov et al., 1996) (Shchelkunov et al., 1996) (Shchelkunov et al., 1996) (Safronov et al., 1996) (Goebel et al., 1996)
173R <i>B5R</i> B7R	155424 156377	317 317 317 259	35.1	ps/hr protein/ EEV gp42 complement control protein VAC VAR-I (BSH:B6R) CPX D17L 186 matches to complement control protein family	1.6e-232 7.1e-220 2.1e-12 <7.7e-05	312/317 294/316 16/52	98 93 30	(Takahashi-Nishimaki et al., 1991) (Engelstad et al., 1992) (Isaacs et al., 1992) (Goebel et al., 1990) (Shchelkunov et al., 1995) (Safronov et al., 1996)
174R 86R 87R	156474 156995	173 173 65 685	20.2	20.2k protein VAC VAR-BSH (I:B8R) NAD-protein ADP ribosyl- transferase phage T4	1.5e-121 6.0e-40 0.56	173/173 62/65 17/56	100 95 30	(Gocbel et al., 1990) (Shchelkunov et al., 1995) SXBPT4
175R <i>B7R</i>	157033 157566	177 182 184 182	20.7	20.7k protein VAC VAC CSL CPX D12L EF-hand calcium-binding domain	7.8c-129 0.16 0.49	95/108 9/44 8/36	87 20 22	(Goebel et al., 1990) (Goebel et al., 1990) (Safronov et al., 1996)
176R 88R 88R	157621 158301	226 272 266 266 274		31k interferon-gamma receptor (f) VAC VAR-BSH (I:B9R) ECT swinepox C6	3.3e-164 3.0e-153 2.6e-151 3.2e-09	116/123 111/123 110/123 12/31	94 90 89 38	(Upton et al., 1992) (Alcami and Smith, 1995) (Goebel et al., 1990) (Shehelkunov et al., 1995) (Mossman et al., 1995b) (Massung et al., 1993)

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<u>ninal</u> 58458 58676		on:	function / homologies	expect	AA id	(%	references
		8.3	8.3k protein				
		0.5	8.3k protein VAC	3.0c-49	60/60	100	. (0-1)
	240		- Capripox T4 protein	1.2e-09	16/44	36	) (Goebel et al., 1990) M28823
	237		shope fibroma virus	0.0057	15/50	30	F43692
58639	158	17.9	17.9k protein				•
59115		.,.,	VAC	4.7e-110	146/146		
	530		swinepox VC04	0.040	13/42	5 100 30	(Goebel et al., 1990) (Massung et al., 1993)
	689		kelch protein D. melanogaster	0.14	12/54	27	(Xue and Cooley, 1993)
59187	74	8.5	8.5k protein				(Senkevich et al., 1993b)
59411		0.5	VAC	9.2e-43	70/72	0.6	
			177 matches to glu/asn rich	9.26-43	70/73	95	(Goebel et al., 1990)
			proteins				
59478	283	33.3					
60329	283	33.3	protein kluase VAC	1 0- 207	202/002		(Howard and Smith, 1989
	134		VAR-I	1.8e-207 8.7e-26	282/283 31/54	99 57	(Goedel el al., 1990)
	300		VACBIR	1.7c-54	26/53	49	(Shcheikunov et al., 199 (Goebel et al., 1990)
	300		VAR-I BIR	7.7e-53	25/53	47	(Shchelkunov et al., 199
			t20 matches mainly to protein kinase family	<0.34			
	٠.		Zinase family				
60437	116.	13.0	ICE inhibitor / SPI-2 (f1)				(Kotum) and M
50787							(Kotwal and Moss, 1989) (Smith et al., 1989)
	116		VAC				(Ray et al., 1992)
	344		VAR-I (BSH:B12R) CPX crmA	3.0e-72	111/116		(Goebel et al., 1990)
	341		VAC CI2L (SPI-I)	2.7e-69 2.8c-39	105/114 66/100		(Shchelkunov et al., 1995
	353		Ectromelia serpin	2.1c-23	25/34	66 73	(Pickup et al., 1986) (Goebel et al., 1990)
	344		rabbitpox SPI-I	9.2c-23	24/34	70	(Senkevich et al., 1993b)
	357 355		CPX SPI-I	5.5c-22	25/34	73	(Ali et al., [994)
	372		VAR-I B25R (BSH:B21R) CPX serpin-like protein	1.4e-21	25/36	69	(Ali et al., 1994)
	372		135 matches mainly to serpins	1.7c-21 1.7c-36	25/34 25/36	73 69	(Shcheikunov et al., 1995 (Ali et al., 1994)
				⊲0.12	25750	03	(An et at., 1994)
0762 1430	222 222	24.9	ICE inhibitor/SPI-2 (f2)				see above
. 770	345		VAC VAC WR	6.2e-158	218/222	98	(Goebel et al., 1990)
	345		rabbit pox SPI-2	9.4c-156 1.6c-153	215/221	97 05	(Kotwal and Moss, 1989)
	341		CPX crmA	4.5c-148	203/220	95 92	(Ali et al., 1994) (Pickup et al., 1986)
	344		VAR-I (BSH:B12R)	1.5c-146	203/220	92	(Shehelkunov et al., 1995)
			309 matches see above	<1.3c-21			
1506	143	16.7	16.7k protein				45 14 .
	149		VAC	3.6e-105	97/98	98	(Smith and Chan, 1991)
	149		VAR-I(BSH:BI3R)	9.1c-104	95/98	98 96	(Goebel et al., 1990) (Shehelkunov et al. 1995)
	153		VAR-I DIL (BSH:D2L)	8.8e-31	25/52	48	(Shchelkunov et al., 1995 (Shchelkunov et al., 1995
	181 159		VAC C16L/B22R	1.0e-26	25/52	48	(Goebel et al., 1990)
	151		capripox T3A rabbit fibroma T3A	1.4c-17 2.6c-07	17/42 17/44	40	(Gershon and Black, 1989
	190		VAC A52R	0.073	10/28	38 35	(Uplon es al., 1987)
	149		VAC WR K7R	0.21	7/22	31	(Goebel et al., 1990) (Boursnell et al., 1988)
	149 161		VAR-I C4R CPX M6R	0.30	7/22	31	(Shehelkunov et al., 1995)
	101		CFA MOR	0.51	7/22	31	(Safronov et al., 1996)
	326	36.6	interleukin-18 receptor				
1001			(IL-18R)				(Alcami and Smith, 1992)
	326		VAC-WR BISR	2.8e-229	323/326	99	(Spriggs et al., 1992) (Smith et al., 1991)
	326 290		CPX B16	2.3c-217	306/326	93	(Spriggs et al., 1992)
	69		VAC VAR-I (BSH:deleted)	4.4c-202 8.1e-38	287/290	98	(Gocbel et al., 1990)
	296		HS type II IL-1 receptor	1.7c-36	59/68 28/75	86 37	(Shchelkunov et al., 1995) U64094
			271 matches mainly to IL-1	<0.011	_0,.,	31	
			receptors, growth factor				
			receptors and lg family proteins				
			b				
		39.6	39.6k protein				
	340			4.8e-248	335/340	98	(Goebel et al., 1990)
	340		VAR-BSH (I:B18L)	2.7e-241	325/340		(Shchelkunov et al., 1995)
209	574	68.0	68k unk-like protein				•
933	574		VAC	0.0	560/574	97	(Smith et al., 1991)
:	574		VAR-I (BSH:BI6R)		539/574	97	(Goebel et al., 1990) (Sheheikunov et al., 1995)
			100 matches mainly to				(=====================================
			poxvirus ankyrin proteins				
000	234	27.5	surface antigen.				//tada
999 2							(Ueda <i>et al.</i> , 1990) (Symons <i>et al.</i> , 1995)
999 2 703			receptor (f)				(Symons et al., 1995) (Colamonici et al., 1995)
703						93	(Goebel et al., 1990)
703	4.3%		***		111/133	83	(Shehelkunov et al., 1995)
703 1 3				v.0031	13/43	34	(McMahan et al., 1991)
703 1 3							
703 1 3				<0.53	-		
00~	:	353 354	234 27.5 353 354	100 matches mainly to poxvirus ankyrin proteins  234 27.5 surface antigen, IFN-alpha/beta receptor (f)  353 VAC (WR:818R)  354 VAR-I (BSH:B17R)	100 matches mainly to poxvirus ankyrin proteins   40.53	VAR-I (BSH:B16R) 0.0 539/574  100 matches mainly to	VAR-I (BSH:B16R) 0.0 539/574 93 100 matches mainly to c0.53 234 27.5 surface antigen, IFN-alpha/beta receptor (f) 353 VAR-I (BSH:B17R) 1.53-149 111/133 83 569 HS interleukin-I receptor 0.0051 15/43 34

ORF'	START	AAb	kDaʻ	name / (putative)	BLAST*	BLAST	HSS	references
1.61	STOP			function / homologies	expect	AA id	(%)	
	erminal	regio	n:					
B22R	167414	1897		VAR-BSH (I:B26R)	9.9e-23	31/38	81	(Shchelkunov et al., 1995)
189R	167897	188	21.7	21.7k protein				
B22R	168463	181		VAC B22R/C16L	2.9c-111	95/104	0.1	(0.11.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.
DIL		153		VAR-I (BSH:D2L)	1.2e-88	66/71	91	(Goebel et al., 1990)
		149		VAC BISR	7.2e-19	25/52	92	(Shchelkunov et al., 1995)
		159		capripox T3A	8.0e-05	15/45	48	(Gocbel et al., 1990)
		151		VAC C6L	0.25	12/46	33	(Gershon and Black, 1989a
		156		VAR (I:D9L;BSH:D12L)	0.26	12/46	26	(Goebel et al., 1990)
				(***=,=====,	0.20	12/40	26	(Shchelkunov et al., 1995)
190R/ 004L	168531 169232	233	26.9	45k ank-like protein (f2)				
B23R	1	386		VAC (C17L/B23R)	6.2e-159	1104110		_
DIL	1	91		VAR-BSH	9.1e-31	110/110		(Goebel et al., 1990)
	1	669		CPX host range	1.1e-13	46/49	93	(Shcheikunov et al., 1995)
	1	452		VAR-I D6L (BSH:D8L)	1.7e-11	22/50	44	(Spehner et al., 1988)
	1.	574		VAR-I BI9R (BSH: BI6R)	1.2e-05	21/50 22/73	42	(Shchelkunov et al., 1995)
	1	574		VAC BISR (WR: BI7R)	8.6e-05	22/73	30	(Shchelkunov et al., 1995)
	1	634		VAC C9L	0.00011	11/24	30	(Goebel et al., 1990)
	1	585		VAR-I GIR	0.00011	22/74	45	(Kotwal and Moss, 1988a)
	Ī	516		orf virus	0.0088	15/49	29	(Shchelkunov et al., 1995)
	J·	153		VAR-I D7L (BSH:D10L)	0.014		30	(Sullivan et al., 1995b)
191R/ 003L	169309 169617	102	12.1	45k ank-like protein	0.014	12/28	42	(Shchelkunov et al., 1995)
B23R	] .	386		VAC CI7L/B23R	1.3e-39	62/63	98	(Goebel et al., 1990)
192R/	170305	176	19.7	secr. TNF receptor (f)				
002L	170835	355		CPX crmB	£ 1 - 71			(Upton et al., 1991a)
G2R		348		VAR-BSH	5.1e-71 1.0e-66	76/83	91 .	(Hu ei al., 1994)
		326		Myxoma virus T2	4.9e-30	73/83	87	(Shchelkunov et al., 1995)
	•	325		Rabbit fibroma Virus T2	1.8e-28	21/37	56	(Upton et al., 1991a)
		202		CPX C4L	8.7e-15	17/36 30/51	47	(Upton et al., 1987)
325R		346		HS TNF receptor	1.9e-08		58	(Heller et al., 1990)
		259		VAC (CI9L/B2SR)	0.00026	14/26 16/19	53	(Safronov et al., 1996)
		277		human CD40L receptor	0.00020	11/24	84 45	(Goebel et al., 1990)
				30 matches to TNF receptors and surface proteins	<0.39	11724	43	(Stamencovic et al., 1989).
93R/	171267	136	14.9	35k major secr. protein				(Development agency
01L	171677			chemokine receptor (f)				(Patel et al., 1990)
129R		244		VAC (C23L/B29R)	6.0e-57	41/42	0.7	(Graham et al., 1997)
35R		253		VAR-I	8.9c-51	46/49	97	(Goebel et al., 1990)
		246		CPX ORF B	5.6e-49	40/49	93	(Shchelkunov et al., 1995)
		258		SFV T1 protein	2.5e-20	23/42	95	(Hu et al., 1994)
		260		Myxoma virus T1/35kDa	1.5e-14	21/42	54 50	(Upton et al., 1987)
						41142	20	(Graham et al., 1997)

<sup>&</sup>lt;sup>a</sup> Open reading frame coding for at least 65 amino acids (for exceptions see text); minor ORFs located in reverse orientation within large ORFs or ORFs located in the repeat regions of the ITRs (see text) are not listed; the MVA ORFs (boldface), listed consecutively as appearing in the genome, and homologs in the Copenhagen strain (in italics), in the variola strains and in the molluscum contagiosum, are listed in this row. Split ORFs are boxed.

b Number of deduced amino acids (AA) encoded within an ORF.

 $<sup>^{\</sup>rm c}$  Predicted  $M_{\rm r}$  (kDa) for the unmodified protein.

<sup>&</sup>lt;sup>d</sup> The lowest Poisson probability determined by the BLAST search (Altschul *et al.*, 1990). The Expect value of 0.0 indicates a probability of zero that an alignment occurs by chance; low Expect values correspond to high homology and vice versa.

<sup>&</sup>lt;sup>e</sup> Amino acid identity (AA id) of first high-scoring segment pair in the BLASTp protocol.

Amino acid Identity of first high-scoring segment pair (HSS)%.

<sup>9</sup> Homologies based on searching PIR and SWISS-PROT databases (BLASTp nr).

h Duplicated ORFs located in ITRs.

Fragment: complete homologous ORF present in related poxvirus (see reference).

J Variola India (I) or variola Bangladesh (BSH) sequences; in cases where the variola sequences are not identical, the variola strain first appearing in the blast search protocol is listed.

k ank, ankyrin.

<sup>&</sup>lt;sup>1</sup>HS, homo sapiens.

<sup>&</sup>lt;sup>m</sup> MM, Mus musculus.

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